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PART 10

**Flue and Exhaust
Gas Analyses**

**INSTRUMENTS
AND
APPARATUS**

**Supplement to
A S M E**

**PERFORMANCE
TEST
CODES**

THE AMERICAN SOCIETY OF MECHANICAL ENGINEERS
United Engineering Center
345 East 47th Street New York, N.Y. 10017

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FOREWORD

The scope of the work of PTC Committee No. 19 on Instruments and Apparatus is to describe the various types of instruments and methods of measurement likely to be prescribed in any of the ASME Performance Test Codes. Such details as the limits and sources of error, method of calibration, precautions, etc., as will determine their range of application are given.

Only the methods of measurement and instruments, including instructions for their use, specified in the individual test codes are mandatory. Other methods of measurement and instruments, that may be treated in the Supplements on Instruments and Apparatus, shall not be used unless agreeable to all the parties to the test.

This Supplement on Instruments and Apparatus, Part 10 on Flue and Exhaust Gas Analyses, replaces an older one published in 1968.

This Edition was approved by the Supervisory Committee on May 15, 1978. It was approved and adopted by the American National Standards Institute as meeting the criteria for an American National Standard on August 17, 1978. Following rescission of a critical manual technique by the original source (ASTM) the document was re-approved by the Supervisory Committee with an alternate manual technique on December 29, 1980 and re-adopted by the American National Standards Institute on June 18, 1981.

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ASME PERFORMANCE TEST CODES

Supplement on Instruments and Apparatus

Part 10 Flue and Exhaust Gas Analyses

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SECTION 1, INTRODUCTION

1.01 Presented are descriptions of methods, apparatus, and calculations which are used in conjunction with Performance Test Codes to determine quantitatively, the gaseous constituents of exhausts resulting from stationary combustion sources. The gases covered by this PTC Supplement are oxygen, carbon dioxide, carbon monoxide, nitrogen, sulfur dioxide, sulfur trioxide, nitric oxide, nitrogen dioxide, hydrogen sulfide, and hydrocarbons. Stationary combustion sources include steam generators, gas turbines, internal combustion engines, incinerators, etc.

1.02 Many methods are available for measuring the constituents in flue and exhaust gases. This PTC Supplement describes in detail the most commonly used instrumentation and analytical procedures used for flue and exhaust gas analyses. Included are instrumental methods as well as manual (normally, wet chemical) methods. The instrumental methods include instruments used for non-continuous or continuous sampling using extractive samples and in-situ type instruments that require no sampling system. The variety of sources and purposes for testing preclude the selection of any one technique as being universally suitable for measuring a given species. The intent of this PTC Supplement is to allow the persons applying a Performance Test Code a choice in selecting the instrumentation or analytical procedures most applicable to their situation. This supplement is not intended as a guide in the performance of compliance tests for federal, state

or local government emission standards but will prove useful in establishing measurement systems when government standards permit a choice.

1.03 The text contains general information on combustion, exhaust gases, sampling systems, measurement systems, and calculations commonly used for exhaust gas testing. Section 4 lists specific methods of extractive gas analysis for each species covered by the PTC Supplement while the methods are described in detail in Appendix I. Sampling requirements, sampling conditioning, calibration methods, special precautions, etc., are included for each species. Sufficient information is included to enable engineering personnel to plan exhaust tests, select equipment, perform the exhaust gas tests, and make the required calculations.

1.04 This Supplement is not a comprehensive survey of techniques available but is instead a systematic approach allowing the analysis system to be best designed. Instrumental methods are continually improving so that any document referring to specific instruments will suffer relatively rapid obsolescence. Reference should be made to continuously updated documents such as Instrumentation for Environmental Monitoring (Lawrence Berkley Laboratories) for correct information on specific instruments.

SECTION 2, GENERAL

2.01 Combustion

2.01.1 Conventional Combustion

2.01.1.1 Conventional combustion systems use fossil fuel in the solid, liquid, or gaseous form. Air is utilized as the source of oxygen for the oxidation of the combustible constituents in the fuel. For most sources the major combustible constituents are elemental carbon and compounds of hydrogen and carbon. In the ideal combustion process, these are oxidized to form carbon dioxide and water vapor. Small quantities of sulfur present in fossil fuels are also combustible and contribute slightly to the heating value of the fuel.

2.01.1.2 Theoretical air is the quantity of air necessary to supply enough oxygen to convert all the carbon, hydrogen, and sulfur in a fuel to carbon dioxide, water and sulfur dioxide. Complete combustion of fuel with no air in excess of the amount theoretically required is an ideal condition. Excess air over the theoretical air is required for satisfactory combustion of the fuel in a reasonable space and time. By the analysis of the exhaust gases, one can determine how well the combustion approaches theoretical stoichiometric conditions.

2.01.1.3 The major gaseous constituents of flue and exhaust streams from normal combustion systems are carbon dioxide, oxygen, nitrogen, and water vapor. In excess air calculations, some or all of these gases must be determined with an accuracy not exceeding ± 2 percent by volume. Equations for determining the excess air, molecular weight, and weight of dry flue gases per pound of fuel are included in Appendix 3.

2.01.1.4 Figure 15 in the Appendix (Par. 6.03.6.4) shows the relationship between oxygen, carbon dioxide, carbon monoxide, nitrogen, and excess air for various fuels of known composition. The oxygen analyzer is an important instrument for combustion control and test purposes because the relationship between oxygen in the flue gases and excess air can be determined from the fuel analysis.

2.01.1.5 The exhaust gases of the combustion process are a major source of heat loss. The sensible heat loss (see PTC 4.1) in the exhaust gases can be determined with acceptable accuracy from a careful measurement of the exhaust gases and their temperatures. It is essential to measure the temperature of these gases to determine the efficiency of the unit.

2.01.1.6 The most important of the trace species found in exhaust gases from normal combustion processes are carbon monoxide, hydrocarbons, nitric oxide, nitrogen dioxide, sulfur dioxide, sulfur trioxide and hydrogen sulfide. When found in exhaust gases, these normally have relatively low concentrations compared to the major gaseous constituents. Instruments used to measure these low concentrations must be accurate in the parts per million range.

2.01.1.7 A complete fuel analysis is required for emission tests. The water vapor, sulfur dioxide, and other components in the exhaust gases can then be calculated for systems without exhaust gas scrubbers. Sulfur trioxide can be estimated based on experience. The moisture in the exhaust gas is the sum of the free moisture in the fuel moisture from burning hydrogen in the fuel and moisture in the combustion air. Approximately 98 percent of the sulfur is converted to SO_2 and 2 percent to SO_3 in the low excess air combustion processes of boiler plants. The analysis may vary widely in other plants. The fuel analysis is often helpful in estimating other component gases.

2.01.2 Other Combustion

2.01.2.1 Some combustion systems do not use fossil fuel. Others do not use air as the source of oxygen for the oxidation. These include units that burn blast furnace gas, coke oven gas, and waste products from the oil refinery industry as well as refuse incinerators and heat recovery boilers that burn domestic or industrial refuse such as sawdust, bark, black liquor, and cane sugar. Other systems use oxygen or exhaust gases that are high in oxygen for the source of oxygen for the combustion process.

2.01.2.2 The normal relationship between excess air, oxygen, and carbon dioxide varies for these special combustion systems. Equations other than those in the Appendix, relating oxygen to excess air must be developed for these special combustion systems.

2.01.2.3 Special combustion systems often include the same trace species found in normal combustion exhaust as well as species formed by the oxidation of other products in the fuel. For example, a wide variety of gaseous species are found in the exhaust from refuse incinerators that burn domestic and industrial refuse that are not found when firing fossil fuel. A complete fuel analysis should be used to determine the possible types of gases and to estimate their concentration. This PTC Supplement only

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covers methods of determining the pollutants that are found in stationary combustion sources burning fossil fuel. Other gases that may be present in the exhaust from special combustion systems may interfere with some of the test methods covered by this text. The instrument manufacturer and the designer of the combustion systems should be consulted in these cases for possible interferences and proper calibration procedures to be used to determine the extent of the interferences.

2.02 Purpose of Test

2.02.1 There are many purposes for making exhaust gas analyses. Table 1 lists some of the purposes for making flue gas tests on a steam generator. For example, O₂ analysis is commonly used to monitor and control combustion processes to obtain safe and efficient operation.

2.02.2 The trace constituents are measured to determine compliance with federal, state, and local regulations and for the monitoring and control of pollution abatement equipment. When compliance with government standards is the purpose of the tests, the testing guidelines published by the federal, state, or local authority must be followed unless a variance can be obtained.

2.03 Measurement Systems

2.03.1 Within the last few years the frequency of exhaust gas analyses has increased in most industries to control various exhaust constituents and unit efficiency. Many manufacturers supply gas analysis equipment using various techniques. The techniques can be grouped into three general areas; i.e., extractive, in-situ, and remote.

2.03.2 The extractive approach has been the most utilized for systems measuring gaseous constituents from stationary sources. This approach requires a sample probe, transport line, conditioning system, and analyzers. There are potential problems in the sampling aspects as well as the measurement.

2.03.3 The spectrometric in-situ approach requires no sample conditioning. The concentrations are determined by the effects on a light (or other energy) source as it passes through the exhaust before impingement on a sensor. The source and sensor are both mounted directly on the stack. They can either be across the stack from each other or colocated on one side with a reflector on the other side of the stack. Although there are no requirements for

TABLE 1
Flue Gas Tests for Steam Generators

Purpose of Test	Constituents	Probe Location
1. Air Infiltration	O ₂ or CO ₂	Furnace and Boiler Outlet
2. Gas Stratification	O ₂ or CO ₂	Furnace, Boiler Outlet, Duct, Stack
3. Air Heater Leakage	O ₂ or CO ₂	Gas Entering and Leaving Air Heater
4. Unit Efficiency	O ₂ , CO ₂ , CO, or HC	Gas at Outlet of Unit (See PTC 4.1)
5. Excess Air	O ₂ or CO ₂	Economizer or Air Heater Outlet
6. Pollution Determination	SO ₂ , SO ₃ , NO, NO ₂ , CO, HC, H ₂ S	Stack or Exhaust Duct
7. Tube Wastage Caused from Reducing Atmosphere	CO, H ₂ S, O ₂	In Furnace at Wastage Area
8. Scrubber Performance	SO ₂	Inlet and Outlet of Scrub- ber
9. Odor Control	H ₂ S	Outlet of Unit
10. Cold End Corrosion (Acid Dew Point)	SO ₃	Air Heater Inlet and Outlet

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sample conditioning with in-stack systems, moisture and particulate in the exhaust can possibly cause interference.

2.03.4 Remote sensing has been developed to enable pollution monitoring from satellite and to allow measurements

of concentration at a distance from the source. This method will require more evaluations in various applications before it can be demonstrated as a viable technique that competes with either the extractive or the in-situ methods.

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SECTION 3, EXTRACTIVE MEASUREMENT SYSTEMS

3.01 There are several considerations that must be made in preparation for an extractive measurement test. They should all be considered or the test may be either under or over designed. These are:

- Purpose of the testing; including the required accuracy.
- The analytical technique to be used.
- Test plan to insure that the data collected fulfills the test requirements.
- The probe design and arrangement.
- The sample transport line and sample conditioning system.

3.02 Sampling Considerations

3.02.1 Test Requirements

3.02.1.1 Before a test can be started, the requirements of the test must be firmly established and suitable equipment chosen. In particular, the accuracy required must be determined and the type of test specified; e.g., whether equipment that will monitor emissions over a period of years is required, whether the test is part of a performance test on the combustion equipment, or whether the test is to prove compliance with regulations.

3.02.1.2 The instrumented or manual analytical method should be chosen to span the entire range of expected gas concentrations. The highest possible concentration should not exceed the range of the method; ideally, the average concentrations should lie at 75 percent of the range. Several calibration points on the analytical system must be made to span the entire range of expected concentrations, as the minimum requirement.

3.02.2 Instrumental or Manual

3.02.2.1 The first choice to be made in an extractive system is between manual (normally wet chemical) and instrument methods. If monitoring over an extended period is specified, then an instrument technique is usually required, and in some compliance testing, manual methods are specified. In most other cases, either method can be used and the choice must be made on the basis of accuracy, convenience, and practicality. In general, instrument methods will be more convenient when many readings are required. Instrumental methods are more expensive and require relatively complicated sample conditioning

and calibrating systems. Manual methods often require only simple apparatus and are inexpensive, but may entail a considerable delay in obtaining results. Manual methods also require careful work by an experienced technician if accurate results are to be obtained. Section 4 on the specific gas analysis should be referenced for selection of a suitable analytical technique.

3.02.3 Test Plan

3.02.3.1 Before the test is started, a detailed test plan should be prepared. This should include shakedown of the instrumentation and sampling system, the calibration procedures and frequency, the number of test runs to be made at each nominally identical condition, and the calculation procedures to be used to convert the raw readings to useful form. The test plan will be influenced by the purpose of the test, the accuracy required, the stability of the combustion process, the exhaust conditions, and the analysis method used.

3.02.3.2 Attention must be given to the method to be used for extracting the sample from the exhaust stream. Since only a very small part of the total flow will be analyzed, it is essential that the gas analyzed is representative of the average exhaust gas. Factors which will influence the accuracy (i.e., the validity) of the sampling system are the design of the sampling probe and the uniformity of the gas composition, velocity and temperature over the duct cross section at the sampling point.

3.02.3.3 Unless prior experience indicates that there is no stratification at the sampling position, it will be necessary to do preliminary testing to measure the velocity, temperature and composition profiles across the sample plane. A sampling procedure can then be selected to give the required accuracy. If the stack has significant stratification that would cause large errors with single point sampling, it may be necessary to traverse the stack and take samples at many points. In some cases, it is possible to design a multi-hole probe that will give acceptable accuracy (see Par. 6.06). If there is little stratification, a single reading from a single-hole probe may be adequate.

3.02.3.4 The stability of the stack conditions with time must be considered before choosing the analysis method and equipment. If conditions vary appreciably, it may be necessary to average results over a period of time. Such averaging will be much easier with instrument systems than with wet chemical methods.

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3.02.3.5 Care must be taken that variations in velocity, temperature, or composition are not caused by changes in the operating conditions occurring while the traverse is being made. A recognized characteristic of the process such as fuel flow or the air flow to fan speed ratio should be monitored during the traverse to verify stable operation. If there is any chance of conditions in the stack changing with time, the readings should be repeated until consistent results are obtained. It may be preferable to use multi-point probes rather than traversing. Otherwise, if conditions continue to change with time, consistent results (i.e., one value) will not be obtained, but an operating range must be established.

3.02.4 Sample Location

3.02.4.1 In many cases, the exhaust ducting system is long, and careful consideration must be given to selecting the sample location. The following should be considered in making this selection:

3.02.4.2 In some cases, the location of the measurement point is determined by the purpose of the test, e.g., measurement of exhaust emissions must be taken after any emission control device while combustion efficiency tests require a location closer to the point of combustion. The concentrations of several gases will vary as the exhaust passes through "scrubbers," and will need to be considered when designing installations for continuous monitoring.

3.02.4.3 The composition of the gas may change with distance along the exhaust system, e.g., NO in the presence of oxygen will change to NO₂ and any heavy hydrocarbons may condense out of the gas stream.

3.02.4.4 If the exhaust stream is cooled sufficiently for water to condense, some components, e.g., NO₂, and SO₂ will be dissolved out of the gas stream into the water.

3.02.4.5 Unless two or more streams are combined, the stratification of gas will be reduced as it passes through the exhaust system, thus allowing a more simple sampling system to be used.

3.02.4.6 Measurements should not be taken within three diameters of the stack exit to avoid the possibility of any ambient air being aspirated into the exhaust and becoming included in the sample. Oxygen readings across the duct should verify that the air is not being drawn in.

3.02.4.7 For continuous systems, the apparatus should be located so that it can be readily inspected and maintained.

3.02.4.8 The analysis should be made in a run of straight uniform ducting so that the velocity profile is as uniform as possible. This reduces the difficulty of obtaining a representative sample.

3.02.4.9 The velocity will be acceptably uniform if the sampling plane is at least eight diameters downstream and two diameters upstream of the nearest disturbance (bend, etc.).

3.02.5 Preliminary Test to Determine Number of Sampling Points

3.02.5.1 When there is no prior knowledge of the homogeneity of the gas concentrations across the sample plane, a preliminary test is necessary to determine the number of sampling positions.

3.02.5.2 It is first necessary to divide the duct cross section into small areas such that the conditions may be assumed to be uniform across each small area. The only requirements in subdividing rectangular ducts are that each of the areas be equal and the proper size, approximately square in shape, and that the sample be extracted at the centroid of each area, as shown in Fig. 1. Sample points in circular ducts are located along diameters normal to each other according to the equation given in Fig. 2.

3.02.5.3 The number of sampling points required in a given duct, may be estimated from Fig. 3. For rectangular ducts, the equivalent diameter is used.

$$\text{Equivalent Diameter} = 2 \left(\frac{\text{length} \times \text{width}}{\text{length} + \text{width}} \right)$$

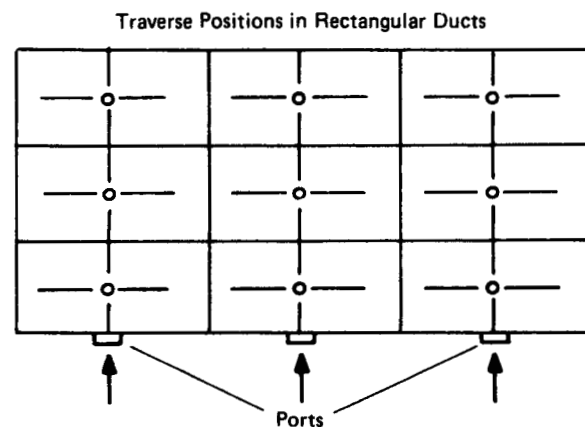
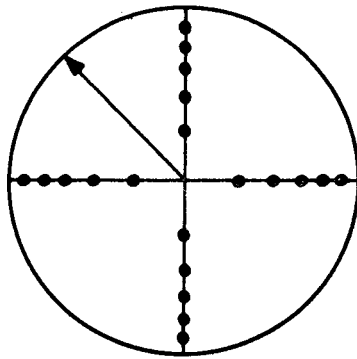


FIG. 1

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Cross Section of
Circular Gas Passage

NOTE: indicates location of
sample point

$$r_n = R \sqrt{\frac{2N_a - 1}{N_T}}$$

r_n = distance from sampling point
to center of pipe

R = radius of pipe

N_a = no. of sampling points counted
from center as zero

N_T = total no. of sampling points on
A diameter

FIG. 2 METHOD OF SUBDIVIDING CIRCULAR SAMPLE PLANE

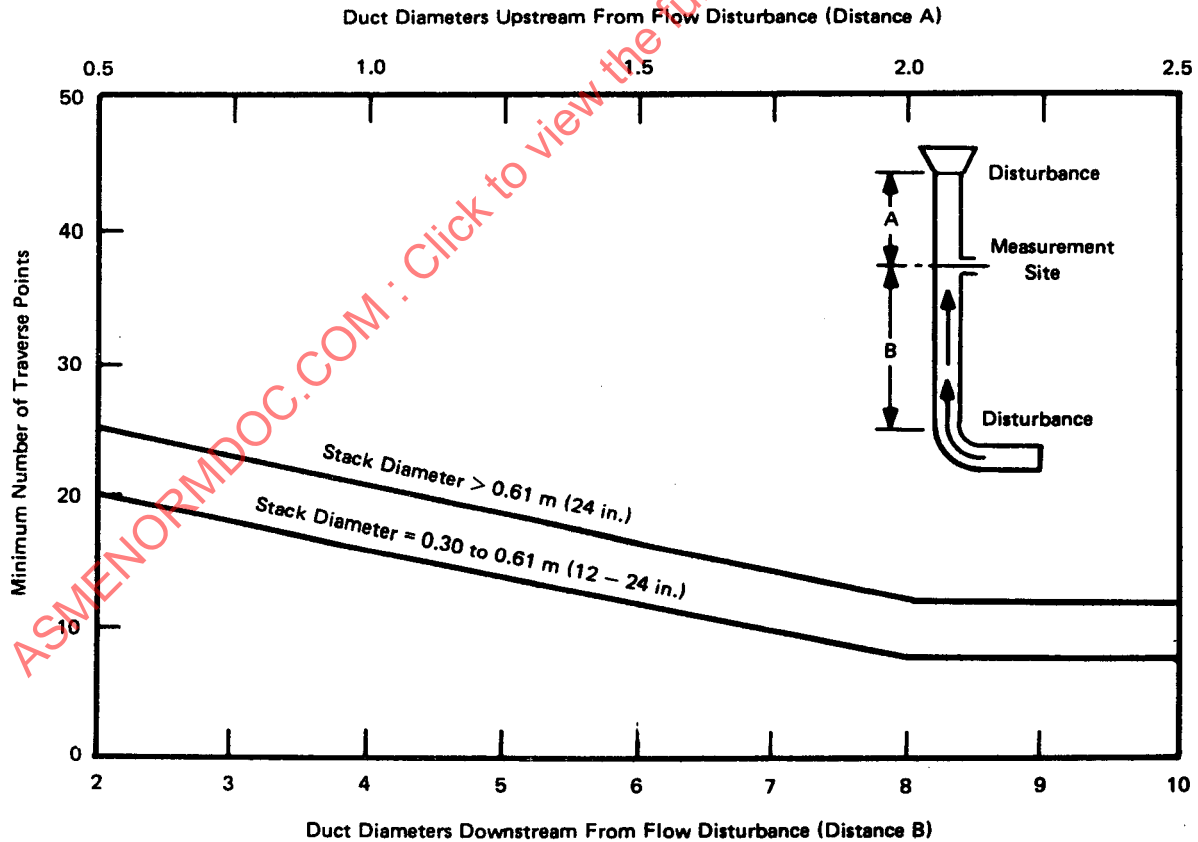


FIG. 3 DUCT DIAMETERS UPSTREAM FROM FLOW-DISTURBANCE (DISTANCE A)

3.02.5.4 Figure 3 will often lead to a conservatively high number of sampling points. This method is the most accurate approach to sampling situations where non-homogeneous flow is possible. A typical test would measure the gas velocity, temperature, and composition at each of the sampling locations given by Figs. 2 and 3. Typical instrumentation for velocity and temperature measurement would be a pitot tube and thermocouple. If there is potential for reverse flow in the duct, a three dimensional probe must be used to determine the net mass flux. If significant gradients are suspected (such as due to a large boundary layer) the suspect areas of the cross section should be further sub-divided. If at all possible, these areas should be avoided.

3.02.5.5 To measure stratification, it is only necessary to measure a single gas component, and this is best done with a continuous instrument. An oxygen or carbon dioxide analyzer would be a good choice, depending on the excess air of the exhaust — the one in higher concentration is used.

3.02.5.6 Once the stratification is known, the minimum number of sampling points required to achieve a desired precision can be calculated. If the preliminary test indicates that the maximum and minimum concentrations obtained were within the required precision, a single point is adequate. If the maximum/minimum concentration variation exceeds the required precision, the mass flow through each small area as determined by the temperature and velocity measurements must be used to calculate the number of points required as outlined in the Appendix (Par. 6.05).

3.02.5.7 The sample analyzed by the instrument should have the same composition as the mass flow average across the sample plane. That is, if the concentration of the constituent in the sample is multiplied by the total exhaust mass flow rate, the result will be total mass flow rate of the constituent.

3.02.5.8 The accuracy given by using a single probe or other sampling system can be calculated from the above information.

3.03 Extractive Systems

3.03.1 Introduction

3.03.1.1 An extractive sampling system is one that removes a portion of the gas inside a flue and delivers it to a chemical or instrumental analyzer. The extractive system must perform three basic functions:

3.03.1.2 (1) Extraction — A portion of the flue gas

that is representative of the sampling plane, must be extracted using a probe or configuration of several probes inserted into the gas stream, as described in par. 3.02.5.

3.03.1.3 (2) Transport — The gas sample is normally drawn through the probe by a source of vacuum and led to the analytical device without chemically altering the species of interest.

3.03.1.4 (3) Conditioning — All components in the exhaust gas sample must be removed which might interfere with the analysis of the species of interest. The state (temperature, pressure, humidity) of the sample must also be suitable for each analytical technique.

3.03.1.5 From the basic hand-held Orsat analyzer to the permanent continuous multi-component stack gas monitors, every extractive system performs these three functions. The degree to which each function is performed is dependent upon the method of analysis and the sensitivity required. The Orsat, for example, is commonly used to measure percentage concentrations of CO, CO₂, and O₂. It uses chemical absorption as a method of analysis. Since it is used near the stack and the reactivities of these gases are low, almost any type of sample line tubing which is free of leaks can be used without interfering with the accuracy. Stack particulate matter is often removed by a cotton or glass wool plug in the sample line, and the probe is normally an unheated stainless or carbon steel tube.

3.03.1.6 Each extractive system must be designed to meet the requirements of both the analytical method used and the specific source tested. Two broad categories are whether the system is to be continuous or intermittent, and whether the analytical method is to be wet chemical or instrumental. Each of these factors requires different hardware to carry out the functions of sample extraction, transport, and conditioning.

3.03.1.7 Calibration is a fourth function of the extractive system which is not necessary for its operation, but determines if the other three functions are satisfactory. The system must be capable of being calibrated through the probe with known concentrations of the gaseous species of interest combined with concentrations of interference gases present in the flue for which the system was designed. The calibrations should include passing the gas through any in-stack filters when they are present. This should include all reactive combinations of gases which might be present in the stack. For example, if SO₂ is being measured, calibration should include SO₂, O₂ and water to determine the amount of SO₂ removed by the sampling system. Such a test need not be performed more than once

ASME PERFORMANCE TEST CODES

on a system if indeed at all, but the system designer must be aware that the data accuracy cannot be assured unless this ultimate test is performed. Routine calibration must be accomplished in the analytical device itself by calibration gases, optical filters, or known solutions. However, where reactive gases are measured, where high sensitivities are required, or where severe stack conditions exist, periodic through-the-probe calibrations are mandatory to insure valid measurements.

3.03.2 Probe

3.03.2.1 The selection of a suitable probe design requires considering three factors: 1) the chemical and physical properties of the gases to be tested, 2) the homogeneity across the sampling plane and 3) flue conditions at the sampling plane.

3.03.2.2 The probe must be capable of withstanding the physical and corrosive environment of the sample plane. In most cases, this requires that it be fabricated from metal. Where the sample gas may react on the metal

surface, glass or ceramic liners are required, but the prime concern is strength.

3.03.2.3 Table 2 shows the minimum reactive material required for sampling each species. Where several species are being measured with the same probe the gas with the most severe requirements dictates the probe material.

3.03.2.4 The internal diameter of the probe should be designed to maintain a high (i.e., turbulent, with a Reynolds number greater than 2300) sample transport velocity throughout. This size should be the same or slightly smaller than the sample line. There should be no dead volumes such as might be created through the use of oversized pipe fittings. If the configuration requires several probes brought together in a manifold, the probes should be proportionally sized or fitted with valves to maintain proper velocity in each.

3.03.2.5 The design of the probe itself is determined by standard engineering practices. Often, it can be as simple as a clear carbon steel or stainless steel pipe. If

TABLE 2
Minimum Reactive Probe Materials to Avoid Sample Reaction

Gas	Temperature At Sample Plane		
	Ambient—750°F	750-1050°F	1050°F and Above
Oxygen	Carbon Steel	Stainless Steel*	Cooled Stainless* below 1200°F
CO	Carbon Steel	Cooled Stainless Steel* 350-900°F	Cooled Stainless* below 900°F
CO ₂	Carbon Steel	Stainless Steel* 350-1050	Cooled Stainless* below 1200°F
NO	Glass or Stainless	Glass or Stainless 350-1050	Quartz or Cooled Stainless below 1200°F
NO ₂	Glass or Stainless Maintained at 350°F	Glass or Stainless at 350°F	Quartz or Cooled Stainless at 350°F
SO ₂	Glass or Stainless Maintained above 350°F	Glass or Cooled Stainless 350-750°F	Quartz or Cooled Stainless 350-750°F
SO ₃	Glass or Stainless Maintained above 350°F	Glass or Stainless 350-750°F	Quartz or Cooled Stainless 350-750°F
HC	Carbon Steel Maintained at 375°F	Stainless Steel* Maintained at 375°F	Cooled Stainless Maintained at 375°F
H ₂ S	Carbon Steel 350-750°F	Stainless Steel* 350-1050°F	Cooled Stainless* 350-1200°F

*Maximum chemical resistance shown by 20 Cb stainless, followed by 316 and then 304.

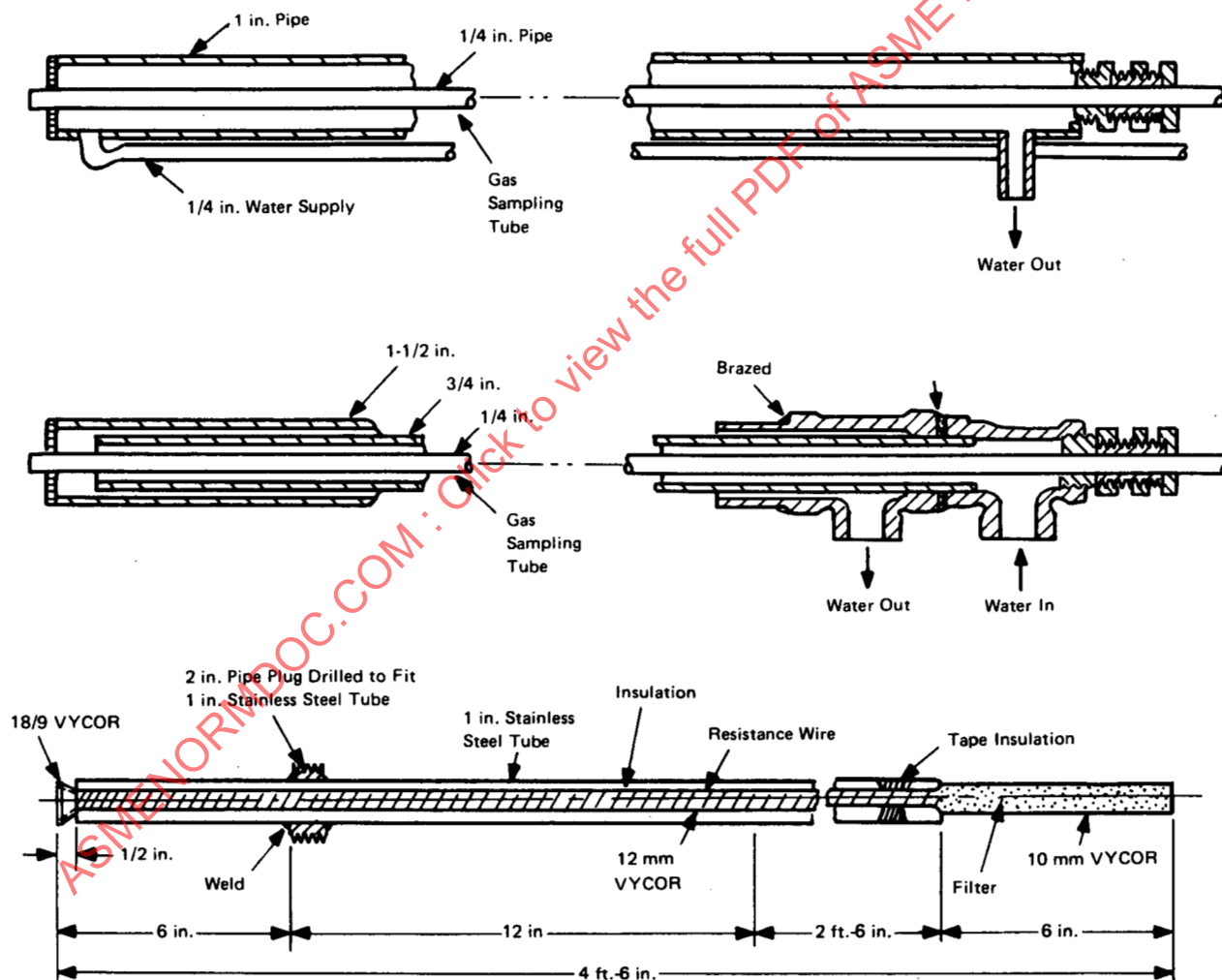
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heating is required, a larger carbon steel jacket is often used to protect the electrical windings. Brazed or welded stainless jackets are used where water, glycol, air, or even steam cooling must be provided to the probe. Where glass or quartz heated liners are necessary, a steel jacket must provide support, protection and allow for thermal expansion between the probe and the liner. Figure 4 shows examples of several typical probe designs.

3.03.2.6 Permanent sampling systems, where wide concentration variations exist, require the use of several separate probes. These can be connected to a manifold or they can be run separately to the analytical device for separate analyses. Figure 5 shows an example of a permanent sampling system installed across a duct. Where

wide variations in flow exist, it may be necessary to adjust the sampling rate at each sample point to be in proportion with the flow at that point to insure that the concentration in the manifold will be representative of the sample plane. Figure 6 shows an example of a system accomplishing this through the use of calibrated orifices. A preliminary traverse is required to determine the flow at each sample point. The sampling location must be chosen to insure that the flows will be approximately proportional to the initially determined flow. This system includes practical means for routine calibration through the sample line.

3.03.2.7 If a single analysis is to be run on integrated (bag) samples from the sample plane and there are wide



NOTE: Material of construction will depend on exhaust temperature and reactivity of gas species.

FIG. 4 VARIOUS TEMPERATURE CONTROLLED PROBE DESIGNS

variations in mass flow across the sample plane as well as concentration, then it is necessary to sample the centroids of equal areas in proportion to their mass flows. In this case, a pitot tube must be combined with the sample probe and the sample taken at a rate proportional to the mass flow at each point.

3.03.2.8 Integrated bags are used where time average sampling is required or for off-site, high-sensitivity analyses such as gas chromatography. Figure 7 shows a large integrated bag sampler. This design draws the sample into the bag by evacuating the container holding the bag. An advantage of this system is that the sample does not pass through the pump. The recommended material for integrated bags is one which does not react with low concentrations of common hydrocarbons or acid gases.

3.03.2.9 Only gaseous constituents of the exhaust gas are considered in this supplement and it is not necessary to sample isokinetically. That is, the velocity at which gas is drawn into the sample probe need not match the velocity at which gas is flowing in the duct because all the gas species will be sampled without preference due to inertial effects.

3.03.3 Sample Transport

Sample transport is the process of moving the sample from the probe to the analyzer. Sample transport requires a pump or vacuum source and tubing through which to move the sample.

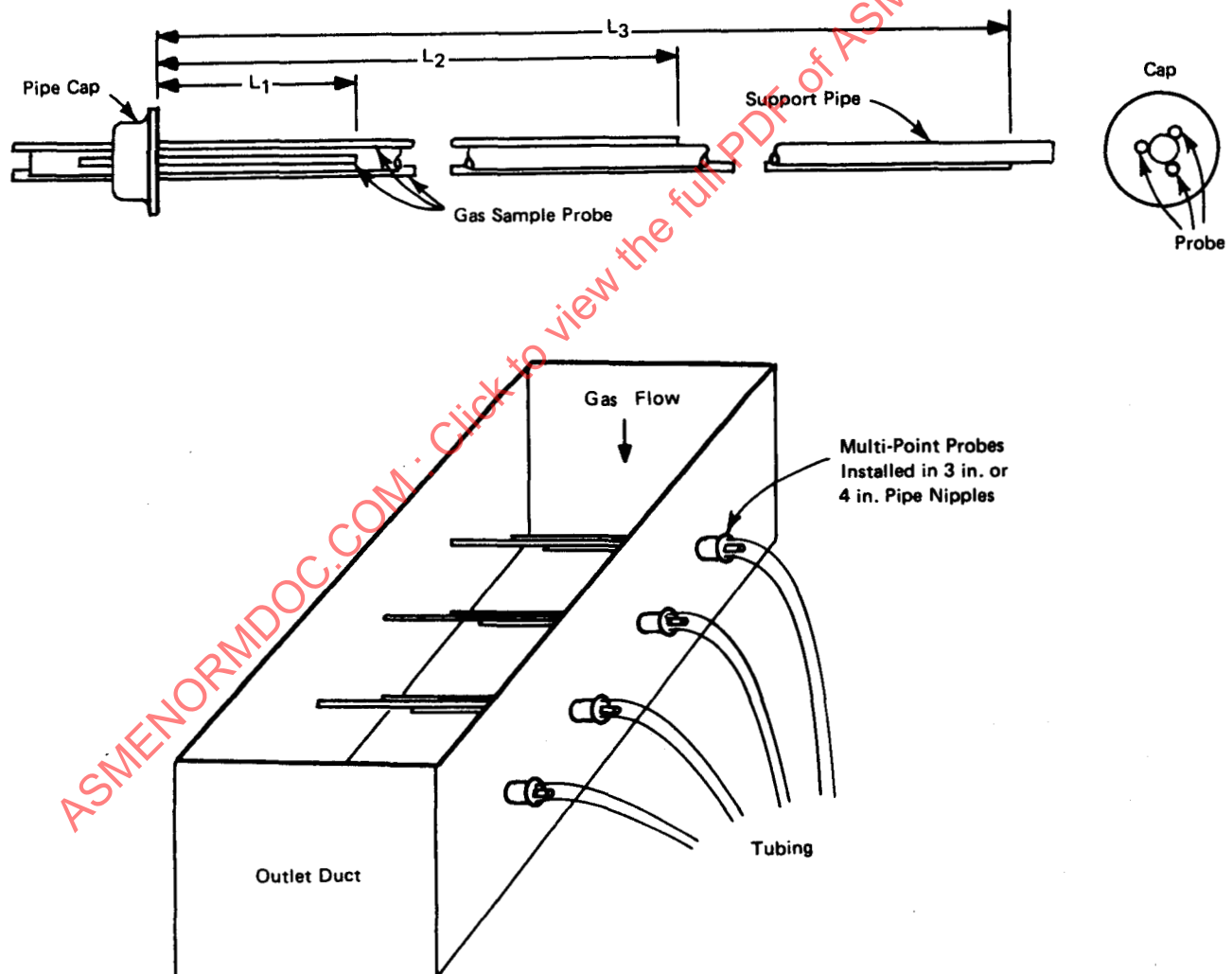


FIG. 5 EXAMPLES OF MULTI-POINT PROBES AND PROBE CONFIGURATIONS.

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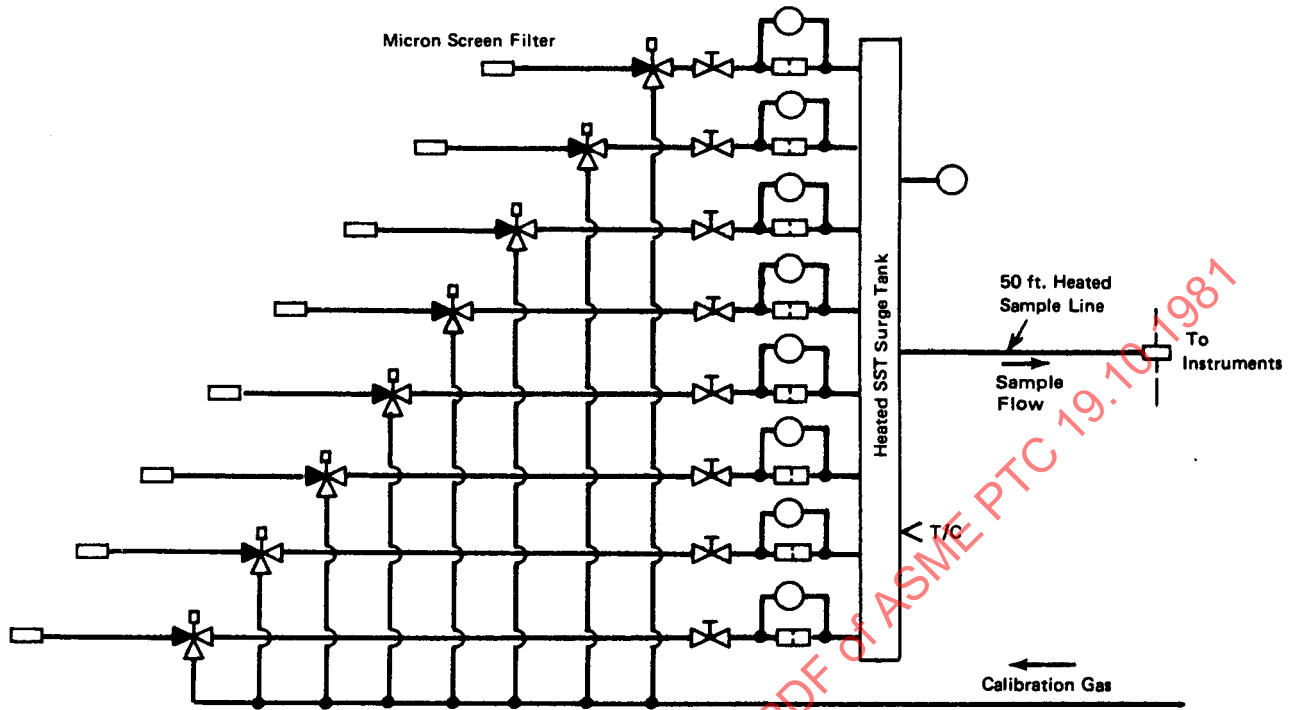


FIG. 6 MULTI-PROBE CONFIGURATION WITH IN-STOCK FILTERS, CONTROLLABLE FLOWS, AND-CALIBRATION SYSTEM

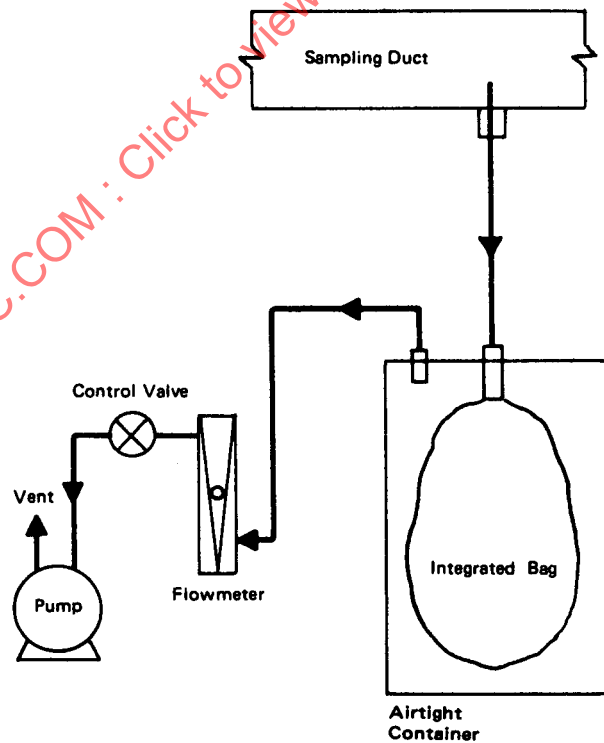


FIG. 7 INTEGRATED GAS SAMPLER

3.03.3.1 Pumps

3.03.3.1.1 Most analytical devices are pressure sensitive. The normal means of allowing for this is to vent the analyzer to atmosphere which requires a pressurized sample stream. Therefore, a pressure-vacuum pump must normally be inserted in the sample line between the probe and the analyzer. Since the gas temperature at the pump will always be less than 400°F and the residence time very short, there is less concern of chemical interactions with the pump materials than with the sample line materials. Certain guidelines must be followed. Oil-lubricated vane or piston pumps should never be used due to the solubility or reactivity of most trace constituents in lubricating oil. Rubber diaphragm pumps are usually adequate except for the measurement of hydrocarbons or the more reactive gases. Teflon diaphragm, stainless bellows and carbon vane pumps are best for all gases. Air and water aspirators are useful for drawing sample through the probe into a manifold outside the stack in locations where electrical power is not easily available or where fire hazards exist. A sample line is then run from the manifold to the analysis area where the electrical pump and instruments are housed. The pump should be located after the moisture and particulate removal devices to increase pump reliability.

3.03.3.1.2 In continuous monitoring systems, it will often be useful to operate the aspirator or pump at a higher capacity than is required for the analytical system. This serves several functions. It produces a short sample transport time between probe and analyzer where long sample lines are required. Also, it helps minimize particulate or moisture drop-out and blockages from occurring in the sample transport system. The excess sample gas is externally vented at the analyzer area. Care must be taken that the vented gas does not interfere with the equipment or pollute the work area.

3.03.3.1.3 The maximum sample transport time between probe and analyzer can be as long as several minutes for non-reactive gases. Very short transport times of 2 seconds have been proposed for sampling reactive gases such as NO. NO reacts slowly with oxygen at ambient temperatures to form NO₂. In heated Teflon sampling lines, the reaction rate is sufficiently slow that with the NO concentrations normally encountered, little oxidation will occur as long as the transport time is less than 10 seconds. Since the equilibrium of this reaction is strongly biased towards NO₂, any dirt in the sample line will act as a catalyst and increase the reaction rate. In all cases it is recommended that the sample transport velocity be maintained high throughout so that the flow is turbulent: i. e., with a Reynolds number greater than 2300.

3.03.3.1.4 Sample flow rates 10 to 100 times that necessary for analysis are common in continuous systems. Preliminary filters and probe and line heaters must be sized accordingly. The valving used to vent the bypass flow must be designed to provide a constant low pressure flow to the analyzers. This must be maintained at all times since most instruments are sensitive to changes in sample pressure. Depending on the instrument, temperature would also be significant but the pumps will normally not influence this. The sample flow rate is not a sensitive parameter for most instruments but should be kept relatively constant. Manufacturer specifications should be followed.

3.03.3.2 Sample Line

3.03.3.2.1 The materials and design of the sample lines are of primary importance in designing a sampling system because of the high surface area to volume ratio of small diameter tubing. For CO, CO₂, and O₂ measurements, aluminum, nylon, polypropylene, or polyethylene tubing is satisfactory. For all other pollutant, either Teflon or 316 stainless steel tubing must be used. Routine cleaning and leak checking of all sample lines is also very important in any permanent installation.

3.03.3.2.2 The sample lines should be constructed with minimum sharp turns, elbows, or unions. Long, straight or slightly curved runs of tubing are desirable where the analyzers are in a remote location. The primary reason is to reduce as far as possible the number of joints which can develop leaks. Small leaks in sample lines which admit air to the sample are very difficult to detect. A thorough leak check must be included during installation of a sampling system and should be repeated routinely at regular intervals. Areas of high vacuum should be avoided throughout the system to minimize air infiltration.

3.03.3.2.3 For gases which do not condense and are not water soluble, sample line heating is not necessary. The line should be physically oriented to prevent condensed moisture from becoming trapped, i.e., it should continuously decline without loops that could collect moisture from the probe to the moisture trap. Where freezing temperatures are likely, the danger of blockage from frozen condensed moisture exists and some form of heating may still be advisable. Insulation alone will not suffice in freezing temperatures.

3.03.3.2.4 Sample lines for CO₂, NO₂, SO₂, SO₃, and H₂S must be heated above the dew point of the exhaust and for SO₃, above the dew point for H₂SO₄. A temperature of 120°C (250°F), is recommended to prevent water condensation. Muller's curve must be referenced (Fig. 8) to determine the line temperature to prevent

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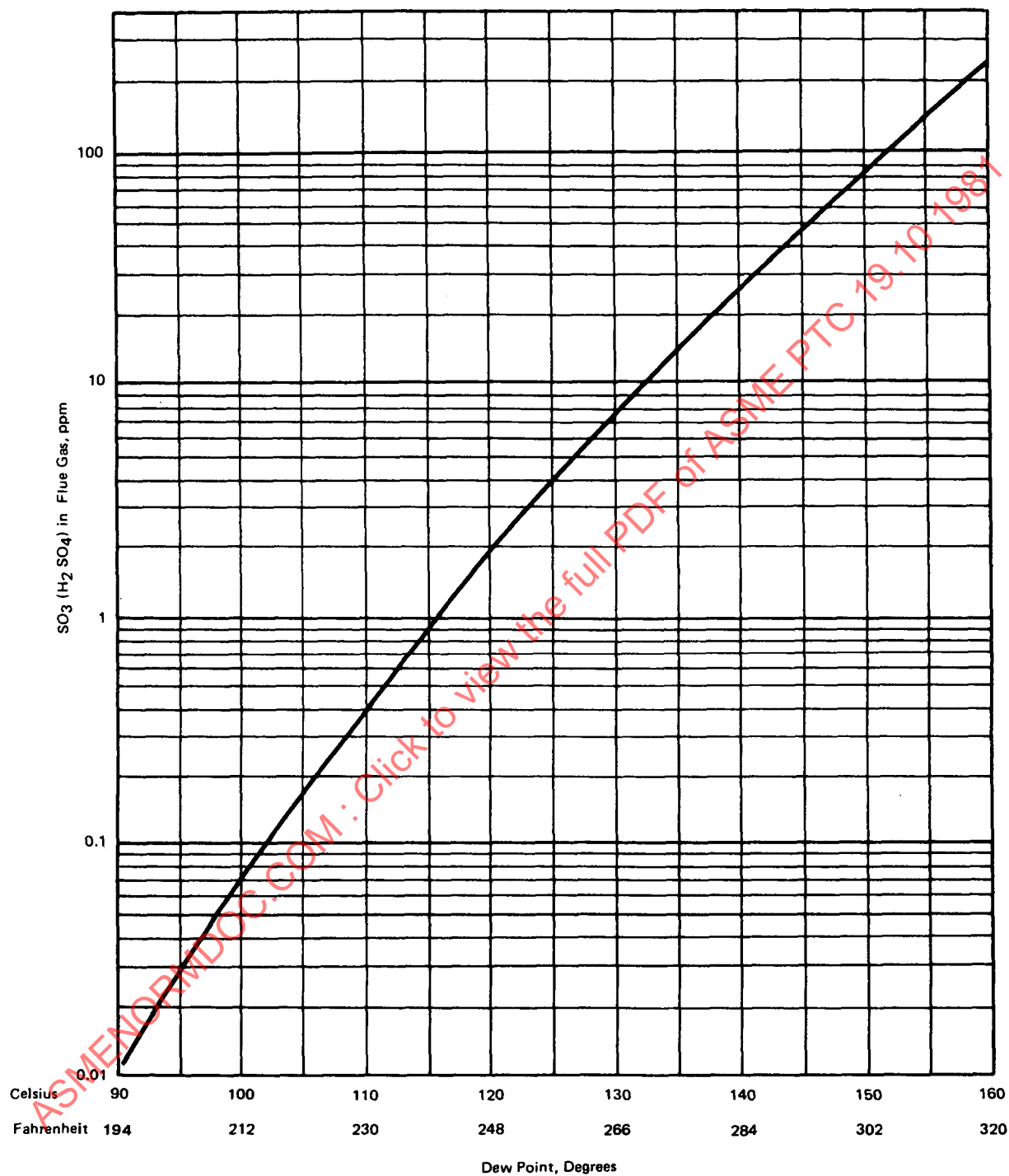


FIG. 8 P. MÜLLER CALCULATED CURVE FOR DEW POINT VS. SULFURIC ACID CONCENTRATION

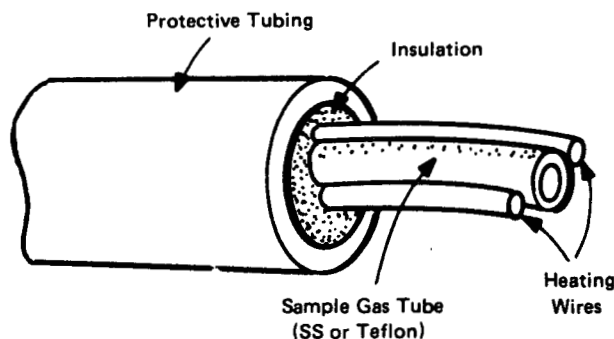


FIG. 9 HEATED TEFLON SAMPLE LINE

H_2SO_4 condensation; generally $150^\circ C$ ($300^\circ F$) is adequate. If high molecular weight hydrocarbons are to be analyzed, the line temperature should be $374^\circ F \pm 9^\circ F$ ($190^\circ C \pm 5^\circ C$) in order to prevent oxidation or condensation from occurring. The heating must be continuous and extend from the probe at the stack wall to the inlet of the analytical device. Due to the high ratio of surface area to volume in small diameter tubing and the low heat capacity of gases, rapid cooling will occur if even a short portion of the sample line is unheated. Special attention should be given to valves and fittings since insulation is inadequate to prevent sample cooling.

3.03.3.2.5 Sample line heaters can be either electrical or steam. An example of a heated Teflon sample line is illustrated in Fig. 9.

3.03.4 Sample Conditioning

3.03.4.1 The process of altering the composition and physical state of the flue gas from the sampling plane before delivering it to the analyzers is termed conditioning. It is a vital part of almost every extractive system if only to the extent of particulate removal and cooling. The separate steps of sample conditioning are: particulate removal, moisture control, temperature control, and removal of interferences.

3.03.4.2 Particulate Removal

3.03.4.2.1 As a general rule, particulate matter should be removed from the sample stream for all instrumented procedures and most wet chemical procedures. The Orsat analyzer is not affected by low concentrations of particulate matter, but often a glass wool or cotton plug is placed in the sample line. A quartz wool plug is required in the peroxide method of SO_2 analysis to remove particulates and acid mist. This is advantageous in most other wet chemical methods where substantial fly ash is encountered.

3.03.4.2.2 A filter is mandatory in continuous instrumental analyzers since even low concentrations of particulate will accumulate in sample lines and probes forming reactive surfaces or flow blockages. Particulates will interfere with pump valving and reduce optical or chemical sensitivities of instruments. The recommended procedure is to use two levels of filtration, a coarse, high flow-rate filter at the probe and a fine replaceable filter immediately before the instruments. This system works well in sampling systems with high bypass flow rates and will assure minimum monitoring system interruptions due to particulate matter.

3.03.4.2.3 A stainless steel filter placed inside the flue on the end of the probe (or probes) can be used if the flue temperatures do not exceed $480^\circ C$ ($900^\circ F$). A cylinder of several layers of fine wire mesh is one commercial version of this filter. A sintered thimble is another alternative. These filters will prove most useful where the majority of

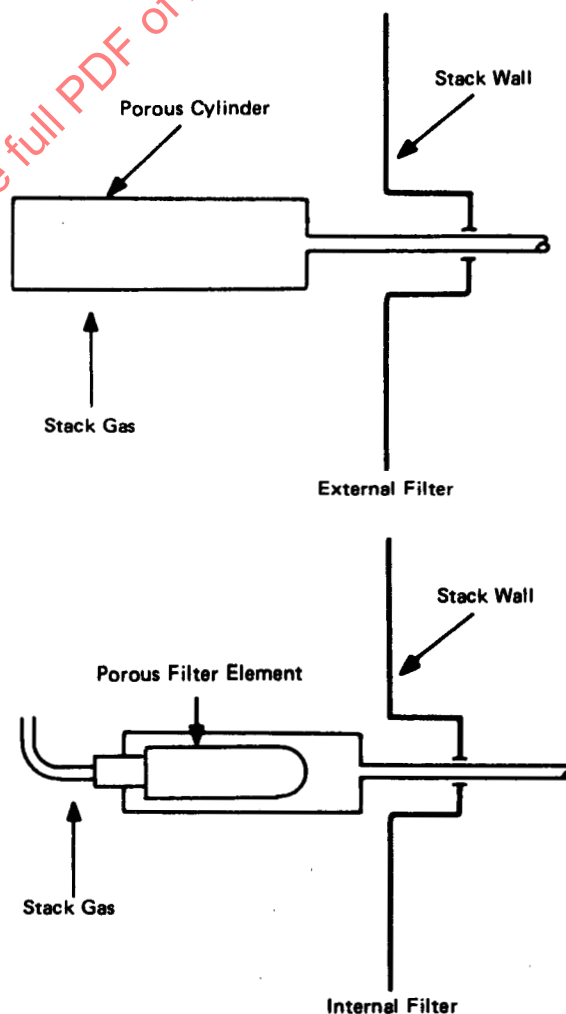


FIG. 10 EXTERNAL AND INTERNAL FILTERS

the particulate in the flue exceeds 20 microns in diameter. The filter porosity should be approximately 60 microns and provision should be made for periodic backflushing of the probe to remove caked-on particulate matter. Table 2 should be consulted to insure material compatibility with the species at the exhaust stream conditions. The purge cycle could be run at the same time as the instrument calibration to minimize data interruption. A metal baffle placed in the flue to protect the instack filter from direct impingement of the gas flow will increase filter life.

3.03.4.2.4 Where higher flue temperature or greater proportions of fine particulate are encountered, a quartz, silicon carbide, or ceramic filter mounted between the probe and the sample line may be used. The filter element is a thimble of no finer porosity than 10 microns which is mounted in an easily accessible, minimum volume holder. Care must be taken to insure that these filters do not absorb or otherwise retain the species being tested. The holder must be heated to the same or higher temperature than the sample line to prevent clogging due to condensation. Since backflushing will not usually be effective in cleaning the entrapped particulate from the filter element, periodic replacement will be required. Figure 10 shows both external and internal filter arrangements. The large surface areas can accumulate material capable of catalyzing reactions involving the species being measured. Calibration will determine the extent of this effect, which cannot be tolerated.

3.03.4.2.5 Reflux filter systems combine primary and secondary filtration techniques to provide a filtration system which can handle large amounts of particulates and reduce particulate size in the conditioned sample stream to less than 1 micron. Figure 11 schematically shows the reflux system. This system incorporates the basic sampling concepts of water or steam washing, but with one major difference; the cleaned sample gas itself is used as the scrubbing medium. This system can also be thought of as a continuous blowback, thus two major primary filter concepts have been incorporated into reflux filtration. The majority of the particulates are rejected and do not enter the probe. A secondary filtration section is required to remove the remaining particulates down to 1 micron or less.

3.03.4.2.6 The operation of the reflux filtration system is as follows: The sample from the stack is drawn through the clean reflux sample stream where dirt particles and aerosols (acid mist) are blown back by the action of high velocity gas molecules. After the sample is scrubbed, it passes through the secondary filter and then is pressurized by a pump. A small portion of the sample is sent to the

dryer (2-3 l/min), while most of the sample (17-20 l/min) is sent back as the clean reflux sample stream. After scrubbing the incoming sample, the reflux stream, along with dirt particles, discharges into the process stream.

3.03.4.2.7 Below 10-20 microns, particulate matter in low concentrations will remain suspended in the sample stream as long as condensation is prevented through heating of the sample line. This finer particulate will still interfere with instrument operation and must be removed in the analysis area. The stainless steel clamshell filter holder containing an easily replaceable fiber glass filter is the best approach. By placing this immediately before the sample pump, the pump and bypass flow valving are protected from particulate matter as well as the instrumentation.

3.03.4.3 Temperature Control

3.03.4.3.1 Analysis systems for hydrocarbons must be maintained at $374 \pm 9^\circ\text{F}$ ($190^\circ\text{C} \pm 5^\circ\text{C}$) from the probe to the flame ionization detector (FID) to prevent the condensation or oxidation of the hydrocarbons. Usually the FID instrument will contain a heated pump so that no other pump is required in the system. A heated fiberglass filter must be placed in the sample line preceding the instrument.

3.03.4.3.2 For sampling, SO_2 , NO , and NO_2 , the sample must be brought to the analysis area at no less than 250°F (121°C). Samples for SO_3 analysis need to be at least 300°F (149°C), see Fig. 8. If instrumental analysis is to be used, these temperatures must be maintained to the instruments. For instruments requiring lower sample temperature the temperature must also be maintained up to the moisture removal device. A heated filter must be inserted in the line. Normally, the pump operating temperature will be sufficient to maintain sample temperature through the pump.

3.03.4.3.3 CO , CO_2 , and H_2S measurements can be made on cool sample streams since these gases are not appreciably moisture soluble at higher concentrations. Condensation knock-out traps must be inserted into the sample line wherever the temperature is lowered. These will usually be part of drying systems as described under Moisture Control.

3.03.4.4 Moisture Control

3.03.4.4.1 Since the moisture in a flue gas stream can vary according to the rate of combustion, the ambient humidity, and the fuel-contained moisture, most gas con-

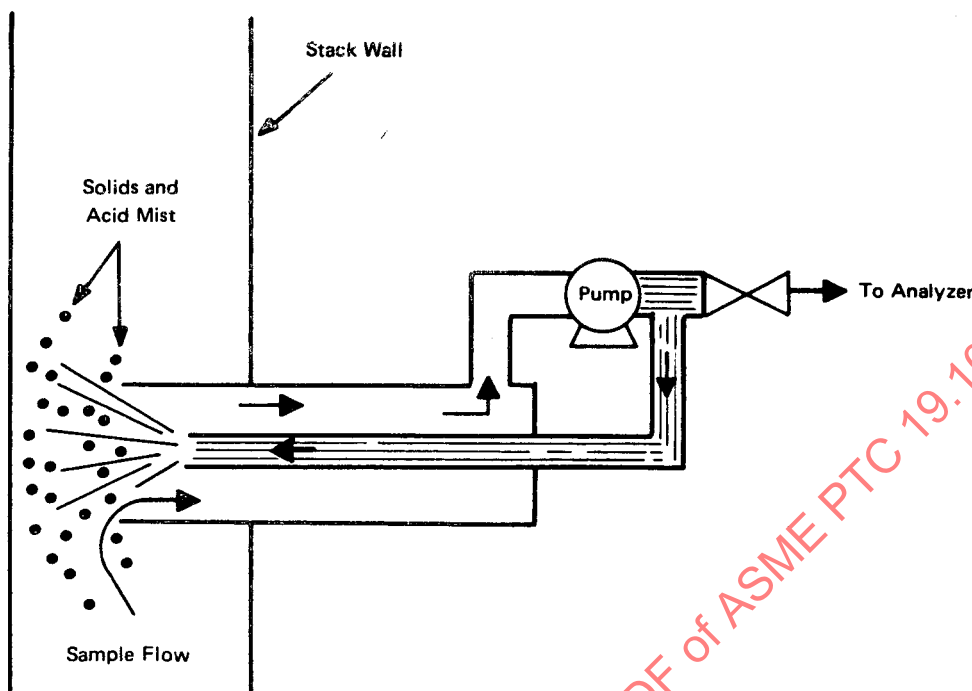


FIG. 11 REFLUX FILTRATION

centrations are reported on either a dry basis or calculated to a constant percent moisture. Ideally, the moisture should not condense before the moisture removal device and it should condense rapidly in this device. In analyzing for CO, CO₂, O₂, and H₂S, it is possible to completely dry the gas stream prior to analysis without appreciably affecting measurements. This is preferable when using non-dispersive infrared (NDIR), gas chromatographic, para-magnetic, and most other instrumental detection systems since the possibility of condensation and consequent reduced instrument performance is eliminated. In the NDIR CO analyzer, moisture interference is also eliminated by complete drying of the sample.

3.03.4.4.2 The air condenser consisting of copper, stainless or glass coil of tubing connected to a liquid trap is often sufficient for the collection of integrated bag samples. These may be used for Orsat analysis or the analysis of non-condensable, non-soluble gases such as methane. Figure 12 shows a schematic of an air-cooled condenser used with an integrated bag sampler. The efficiency of this trap is improved by placing the coil in an ice bath. This arrangement is useful in short-term instrumental monitoring.

3.03.4.4.3 For continuous monitoring, the use of a stainless coil in a refrigerated bath will give long, trouble-free service. It should be designed to minimize contact

between the condensed liquid and the sample stream. Also automatic draining should be provided.

3.03.4.4.4 The permeation dryer also removes virtually all the water in a sample gas stream without affecting the gas concentrations. The wet stream is contacted with an ion-exchange membrane which is permeable to moisture in the gas phase. The other side of the membrane is either evacuated or purged with dry air. The sample line and the dryer must be heated above the dew point since only gas phase moisture can be removed. The permeation dryer has the advantages of not requiring refrigeration or a condensate trap. It has been recommended for systems measuring NO₂ and SO₂ in sources having a low particulate concentration.*

3.03.4.4.5 A less desirable method for controlling condensation is by dilution of the sample with clean dry air using permeable membranes or flow proportionation. These are not recommended because of the possibility of additional error from imprecise calibration.

3.03.4.4.6 Desiccants may be used for drying certain gases such as CO but generally interfere with most other pollutants.

*Walden Research Report — EPA — 650/2-74-089 (October, 1974)
EPA, Research Triangle Park, N.C. 27711

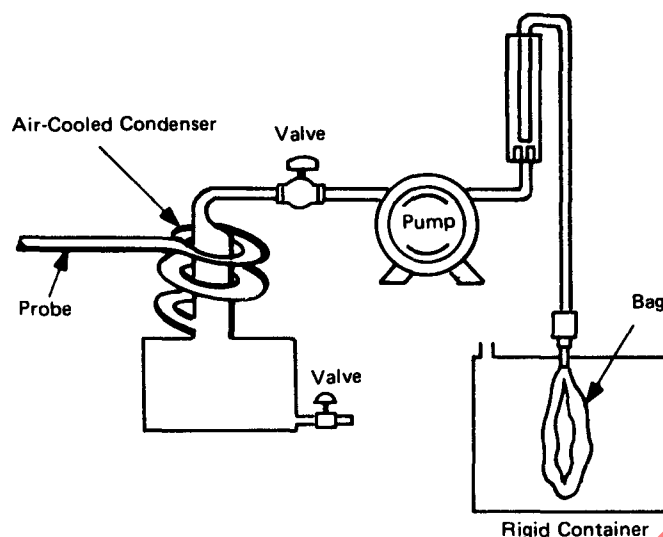


FIG. 12 AIR-COOLED CONDENSER ON INTEGRATED BAG SAMPLER

3.03.4.5 Removal of Interference Gases

3.03.4.5.1 No generalizations can be made concerning interfering gases with the exception of an unalterable rule: Every analytical method or instrument which is used in an extractive measurement system must be examined with respect to its sensitivity to all gases which might reasonably be expected in the flue. Where interferences are found, the sample conditioning system must be designed to remove the interfering gas.

3.03.4.5.2 An example already given was the interference of moisture on CO measurements by NDIR. This is removed by condensation often followed by contact with a drying agent. Electro-chemical membrane instruments are often sensitive to more than one constituent of a flue and require chemical scrubbers or internal electrical

compensation. The latter is valid over a given range and requires adjustment if the interferer concentration changes appreciably.

3.03.4.5.3 Wet chemical analysis methods are also affected by interferences. In the analysis of SO_2 by peroxide oxidation, SO_3 must first be removed since it would be indistinguishable from the oxidized SO_2 . This is accomplished with a preliminary isopropanol scrubber. The isopropanol scrubber is not totally satisfactory in that it removes significant amounts of SO_2 as well; it has been observed to remove 10 percent of the SO_2 at concentrations up to 150 ppm. However, there currently is nothing better available. The definitive procedure for each method must be studied to determine its applicability in the flue gas of interest.

SECTION 4, SPECIFIC GAS ANALYSIS FOR EXTRACTIVE SYSTEMS

4.01 Introduction

This section reviews the most commonly used techniques available for analysis of each species covered by this supplement. Peculiarities of the extractive system as applied to each species are also described here. Further requirements specific to individual analytical techniques are found in the Appendix.

4.02 Oxygen

4.02.1 The measurement of oxygen concentration in flue gas is used to determine the average molecular weight of the flue gas and the excess air above theoretical with which the combustion process takes place. The excess air measurement is needed to determine the gas weights leaving the combustor and the efficiency of the unit. Oxygen is often monitored to control flame stability at low excess air conditions. Oxygen concentration in exhaust gases are in the range of 0 to 21 percent when using atmospheric air for combustion. All determination methods described are applicable to this concentration range.

The sampling system for oxygen requires no special precautions due to its stability in the concentration ranges normally measured.

4.02.2 Manual Methods

The Orsat apparatus is the commonly used manual method for O_2 determination in exhaust gases. A known volume of gas sample is passed through a reagent which absorbs the oxygen. The O_2 content is then determined by the decrease in gas volume. A detailed description and operating procedure of the Orsat are presented in the Appendix (Par. 6.01.1).

4.02.3 Instrumented Methods

The available instruments for O_2 determination operate on the principles of paramagnetic attraction, catalytic combustion, electro-chemical membrane diffusion or diffusion in a zirconia cell. These methods are detailed in the Appendix (Par. 6.02.1-4).

4.03 Carbon Dioxide

4.03.1 Carbon dioxide concentrations range theoretically from 0 to 21 percent (by volume) in flue and exhaust gases when using atmospheric air for combustion. The sampling system for carbon dioxide requires no special precautions due to the stability of CO_2 and the concentration ranges normally measured. Determination methods described

herein are applicable to this concentration range. However, the alkaline scrubber fluids in most power plants equipped with scrubbers will absorb CO_2 quite readily.

4.03.2 Manual Method

The Orsat apparatus is commonly used for manually determining CO_2 in exhaust gases. A measured gas sample is passed through a pipette containing a reagent where the CO_2 is absorbed by contact with the reagent. The CO_2 concentration is determined by the decrease in gas volume after the gas is passed through the CO_2 absorbent. A detailed description and operating procedure of the Orsat are presented in the Appendix (Par. 6.01.1).

4.03.3 Instrumented Methods

The instruments available for CO_2 determination operate on the principle of non-dispersive infrared absorption. This method is described in the Appendix (Par. 6.02.5).

4.04 Carbon Monoxide

4.04.1 Carbon monoxide is a colorless, odorless, tasteless gas, slightly lighter than air. It is produced during incomplete combustion of carbonaceous fuels. Since it is a pollutant gas and represents incomplete combustion, normal combustion systems are adjusted to minimize carbon monoxide concentrations. Measurements in these systems can range from 0 to 500 ppm. In unusual combustors, such as blast furnaces, the CO concentration can range up to 30 percent. Different methods of determining CO concentrations are applicable to different sources because of the wide variation of concentration.

The sampling system for measuring CO must be designed to avoid extremely high probe temperatures which can cause catalytic oxidation of CO.

4.04.2 Manual Methods

The Orsat apparatus may be used for CO measurement in applications where substantial quantities of CO are produced. A typical application would be a coke oven, where there is 5 to 8 percent CO concentration in the flue gas. The Orsat should not be used where less than 1 percent CO is found because the Orsat's minimum sensitivity is 0.5 percent. The Orsat commonly has a 0 to 20 percent range. A measured gas sample is bubbled through a pipette where CO is absorbed by a reagent in the pipette. The CO concentration is determined by the decrease in gas volume after

passing through the CO absorbent. A detailed description and operating instructions of the Orsat are presented in Appendix 1 (Par. 6.01.1).

4.04.3 Instrumented Methods

The instruments available for CO determination operate on the principles of non-dispersive infrared absorption, electrochemical membrane diffusion or gas chromatography with or without a reducing cell and FID. These methods are described in the Appendix (Par. 6.02).

4.05 Nitrogen

4.05.1 Free nitrogen in exhaust gas is not considered a pollutant. 78 percent of the atmosphere is made up of this gas. The free nitrogen content of exhaust gas may contain nitrogen liberated from the fuel as well as unreacted nitrogen contributed by the combustion air. The free nitrogen content of exhaust gas can be determined by subtracting from 100 percent the measured quantities of the other gases. For example, if the pollutants are considered negligible, then the nitrogen can be determined with an Orsat analysis once the O₂ and CO₂ content have been measured.

4.06 Sulfur Oxides

4.06.1.1 Sulfur oxides are emitted as air pollutants from industrial processes such as smelters, sulfuric acid plants and fossil fuel combustion plants. SO₂ is measured to monitor the efficiency of pollutant abatement equipment and is sometimes monitored continuously for compliance with regulations. SO₃ concentration is measured to control cold end corrosion. Stationary combustion sources emit mostly sulfur dioxide with a small amount of sulfur trioxide. The SO₂, which leaves the stack is eventually oxidized to SO₃ in the atmosphere or when it is washed out of the atmosphere by rain. SO₃ readily combines with atmospheric moisture to form H₂SO₄, sulfuric acid. SO₂ emissions range from zero in combustion with natural gas to over 5000 ppm in lead smelters. The methods described herein are applicable to this range of concentrations.

4.06.1.2 All methods for the determination of sulfur oxides require a heated sample train to eliminate condensation. SO₂ readily dissolves in water and becomes oxidized to SO₃, reducing precision and accuracy in SO₃ collection. In addition, SO₂ measurement will be effected, but to a lesser degree. When calibrating a sampling system for SO₂ or SO₃, water should be present in the moisture trap and oxygen present in the calibration gas.

4.06.2 Manual Method

The Federal EPA Method 6, which is given in Par. 6.01.4, is recommended for applications requiring only measurements of SO₂. The Brookhaven Controlled Condensations System, given in Par. 6.01.5, is recommended for applications requiring both SO₂ and H₂SO₄ vapor measurements. The sample collection hardware and procedures of the controlled condensation system are similar to those of the Intersociety procedures, but include significant changes that improve the technique's accuracy. (Methods 711 and 712, "Methods of Air Sampling and Analysis," Second Edition (Morris Katz, Editor). Amer. Pub. Health Association, 1977.)

4.06.3 Instrumented Methods

Continuous instrumented methods available for the determination of sulfur dioxide operate on various principles including non-dispersive infrared absorption, electrochemical membrane diffusion, electrolytic titration, chemiluminescence and pulsed UV fluorescence. Flame photometry is used to measure the total sulfur of all gaseous compounds in a gas sample or can be combined with gas chromatography to measure the individual sulfur containing species. These methods are described in detail in the Appendix (Par. 6.02).

4.07 Nitrogen Oxides

4.07.1.1 Nitrogen oxides (NO_x) are generated by the oxidation of atmospheric and fuel-bound nitrogen in most combustion processes. The most important NO_x species in exhaust streams are nitric oxide (NO) and nitrogen dioxide (NO₂). NO is a colorless gas while NO₂ gas has a reddish-brown color. NO is the only oxide of nitrogen present in significant concentrations at flue gas temperature. NO₂ reaches a few parts per million at flue gas temperatures but can increase as the NO oxidizes downstream of the combustion. Occasionally, NO and NO₂ are measured individually, but most often the determination of total nitrogen oxides (NO_x) is adequate. The rate of formation of NO is exponentially dependent on the maximum combustion temperature, increases with the square root of combustion pressure, and is significantly increased by reaction of oxygen with nitrogen bound in the fuel. The primary variable affecting the selection of an analytical method is the concentration of NO_x expected. NO_x emission levels vary considerably, from 30 ppm in water injected gas turbines to 1200 ppm in certain coal-fired power plants.

4.07.1.2 The sampling system for measuring NO_x must remove moisture. However, NO_2 will be removed by all but the best moisture traps. Permeation dryers can be used as a means of moisture removal without NO_2 loss.

4.07.1.3 When calibrating systems for either NO or NO_2 , the system should include water in the moisture trap and oxygen in the calibration gas. Oxygen is needed in the calibration gas because both NO and NO_2 may react with oxygen. An inadequately designed sampling system would not show this interfering reaction if calibration gas did not include oxygen.

4.07.2 Manual Method

The phenol-disulfonic acid (PDS) method or the Saltzman method of analysis can be used to measure total NO_x when differentiation of the various oxides of nitrogen is not required. The ASTM/ANSI D-1608-77 for PDS and the Intersociety Committee Method for Saltzman manual determinations of NO_x are presented in the Appendix (Pars. 6.01.2 and 6.01.3, respectively).

4.07.3 Instrumented Methods

The instruments available for NO_x measurement operate on the principles of chemiluminescence, electro-chemical membrane diffusion, or non-dispersive ultraviolet absorption. The methods are described in the Appendix (Par. 6.02).

4.08 Hydrogen Sulfide

4.08.1 Hydrogen sulfide (H_2S) is a colorless gas which has a characteristic odor of rotten eggs. It is generated by kraft paper mills and the catalytic cracking units and desulfurization plants of petroleum refineries. It is also found in boiler plants operating at too low oxygen concentration. Measurements normally must be made down to the range of a few ppb because of the extremely low odor threshold (15ppb) of hydrogen sulfide.

The sampling system for measuring H_2S must be designed to avoid extremely high probe temperatures which can cause catalytic oxidation of H_2S .

4.08.2 Manual Method

The $\text{Cd}(\text{OH})_2$ method of analysis can be used to measure

H_2S concentrations down to 1 ppb. The Intersociety Committee Method is given in the Appendix (Par. 6.01.6).

4.08.3 Instrumented Methods

Commercially available instruments for the determination of hydrogen sulfide emissions operate on principles of either electrolytic titration, electro-chemical membrane diffusion, gas chromatography with flame photometry or non-dispersive ultraviolet absorption. These methods are described in detail in the Appendix (Par. 6.02).

4.09 Total Hydrocarbons

4.09.1 Hydrocarbons include all compounds of hydrogen and carbon. For the purposes of combustion analysis, other organic compounds which contain oxygen, nitrogen, sulfur and the halogens are also classified as hydrocarbons. In most cases, it is sufficient to combine all these compounds into a single measurement known as total hydrocarbons and calculated as equivalent methane (CH_4). Hydrocarbons are present in flue gas from a combustion source due to incomplete oxidation of the fuel. Methane may also be present from atmospheric methane (approximately 2 ppm) which is not oxidized by the flame. Often methane is subtracted from total hydrocarbon measurements since it does not participate in atmospheric smog-forming reactions. Because of economy and efficiency, all stationary combustion processes must be adjusted for sufficient excess air to minimize hydrocarbon emissions (0-50 ppm). Stationary and mobile internal combustion engines produce higher concentrations of hydrocarbons due to the extremely short residence time of the fuel in the combustion chamber. Levels may approach several thousand ppm.

The sampling system for measuring hydrocarbons must be designed to keep the sample stream between 365 and 383°F (185 and 195°C) to prevent condensation and oxidation. There should be no moisture traps in the sample system.

4.09.2 Instrumented Methods

Commercially available instruments for monitoring hydrocarbon emissions operate on the principles of either non-dispersive infrared absorption or flame ionization detection, or gas chromatography with flame ionization detector. These methods are described in the Appendix (Par. 6.02).

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SECTION 5, IN-SITU AND REMOTE SYSTEMS

5.01 In-Situ Systems

5.01.1 In-situ systems offer both advantages and disadvantages over extractive systems. The in-situ system permits one instrument to measure several exhaust species, requires little maintenance and is not subject to errors from a sampling system. However, as with extractive systems, the user must have a thorough knowledge of the condition of his exhaust gas if the data will be useful. The user must realize that the system will probably have to be dedicated to a single exhaust. Individual instrument manufacturers should be contacted with regard to the specific application.

5.01.2 The principle of the in-situ spectrometric system involves sending energy of a known spectrum across the exhaust to a sensor. The alteration of the spectrum by the exhaust is compared to a standard by the system's logic unit to determine the concentration of one or more species. The alteration normally involves absorption of certain wavelengths by the species of concern but the criteria the system used to determine concentrations could be other than simple absorption; e.g., relative absorption at two or more wavelengths or the slope of the absorption as a function of wavelength at a certain wavelength may determine species concentrations. The complexity of the logic in the systems is related to possible interfering species in the exhaust.

5.01.3 The source and sensor of the system must be located at a stack (or duct) penetration. They may be arranged in one of two configurations: (1) the source on one side of the stack (or duct) and the sensor on the other, oriented so that the energy from the source is collected by the sensor; (2) both the source and sensor co-located at the same stack (or duct) penetration with a reflection device (normally mounted on the opposite wall of the stack (or duct) to return the energy to the sensor.

5.01.4 The extractive system drawing through one or more probes delivers a sample taken from discrete points

in the cross section. However, the in-situ system, with its small beam cross section will observe a different concentration if the specie measured is not homogeneous. The observed concentration for either will equal the true concentration when there are no concentration gradients in the stack (or duct). This error may not be too important in many exhaust systems. The in-situ system is accurate if it is corrected for the gradients and if the gradients do not change. The user should also calibrate the in-situ with an extractive system. If the exhaust has previously been probed using extractive systems, knowledge of the exhaust homogeneity can be used to correct in-situ system readings.

5.01.5 The normal means for calibrating an in-situ system is to place a microsample of the gaseous species being observed in the optical path of the system. The sample is contained in a suitable container, such as glass or quartz. An alternate means of calibration uses optical filters which have been correlated to known concentration. However, some species such as SO_3 are not stable even in the micro-sample capsules, and these microsamples have to be calibrated periodically. For most gas species, the microsamples have proven to be quite reliable.

5.02 Remote Systems

5.02.1 Remote measurement systems were developed to measure atmospheric concentrations. In recent years, several remote techniques have been developed for air pollution monitoring purposes. While remote systems have inherent problems that may keep them from being as accurate as systems dedicated to a single stack, they may have a future because of their qualitative accuracy. One application is tracking a plume with a high constituent concentration back to the source. Other applications are ambient monitoring (e.g., from aircraft or satellite) and screening tests. Remote systems which are currently being used are the following: correlation spectrometer, long path absorption method, passive emission method, Lidar and Raman spectrometers.

SECTION 6, APPENDIX

6.01 Manual Methods of Analysis. In all of the manual methods having origin as ASTM or other agency, the user should verify that he is using the most recent version.

6.01.1 Manually Operated Orsat

6.01.1.1.1 The Orsat apparatus shown in Fig. 13 is a commonly used gas analyzer for determining percentage concentrations of CO_2 , O_2 , and CO . Several types have been developed, alike in principle, but differing in detail. Basically, the percentage of the main constituents in flue gas are determined by successive absorption of carbon dioxide, oxygen, and carbon monoxide from a measured gas sample in reagent-filled pipettes. The decrease in volume caused by each absorption under constant pressure and temperature is a measure of each constituent with the remaining portion of the sample considered to be nitrogen. The results obtained with each sample are given on a dry basis, and the water vapor present in the flue gas requires an independent determination. However, the moisture may be obtained by calculation when required to determine the efficiency of the unit.

6.01.1.1.2 The standard Orsat has three absorption pipettes which may be the type that absorbs gas by either bubbling through the reagent or by contact without bubbling. Each pipette shall have a capacity of not less than 50 cc of reagent. The burette shall have a measuring capacity of 50 cc, and it shall be water-jacketed to eliminate errors due to temperature change in the gas.

6.01.1.2 Standard Procedure

6.01.1.2.0 The procedure for preparing the instrument for sampling and for measuring the sample accurately are covered in detail in the Orsat manufacturer's instruction book supplied with the instrument. These instructions should be carefully read and followed. Briefly, the general procedure is as follows:

6.01.1.2.1 Fill the pipettes and burette with the proper solutions and bring the level to the reference mark on each pipette. Care is required when charging the oxygen pipette to minimize exposure of the solution to air and hence exhausting the solution.

6.01.1.2.2 Run a leakage test to be certain the Orsat is tight. This may be accomplished by filling the burette half full of air, closing all valves to include the capillary manifold in the test, and then noting if the confining liquid

continues to rise or fall when the leveling bottle is elevated or lowered. Leakage must not exceed $\frac{1}{4}$ cc in five minutes.

6.01.1.2.3 Insure that the confining solution in the burette is saturated with gas by allowing the gas sample to rapidly bubble through for a minimum of ten minutes.

6.01.1.2.4 When the above requirements have been met, start with the burette full of solution and draw in approximately 50 cc of gas sample by lowering leveling bottle. Close the gas valve and open the atmospheric valve to expel the sample. Repeat this procedure four or five times or until a representative sample is retained in the burette. At this point care must be taken to assure that a 50 cc sample will be analyzed by discharging the excess to atmosphere, being sure no air is drawn into the burette during the leveling process. The liquid level in the burette must be at the zero mark when the surface in the leveling bottle is brought to the same level.

6.01.1.2.5 Raise the leveling bottle and open valve to the carbon dioxide pipette, forcing the sample into the pipette. Slowly pass sample back and forth three times and measure absorption by bringing the liquid level in the bottle even with the level in burette. Pass sample through the CO_2 pipette again and note any change in level. Repeat absorption until there is no further decrease in volume; then take reading. One such check reading is usually sufficient.

6.01.1.2.6 Follow a similar procedure for the oxygen and carbon monoxide determinations, keeping a record of the volume absorbed in each case. The acid cuprous chloride solution used to absorb carbon monoxide will sometimes evolve gas to the sample and affect the volume. The sample should be passed through the reagents in the carbon dioxide pipette before passing it to the carbon monoxide pipette for measurement.

6.01.1.2.7 Finally, expel the sample to atmosphere by bringing the burette solution up to the reference mark, and the Orsat is then ready for the next sample.

6.01.1.3 Precautions

6.01.1.3.1 The following general precautions must be taken to avoid possible error and to obtain the best possible results:

6.01.1.3.2 The manifold capillary tube connecting

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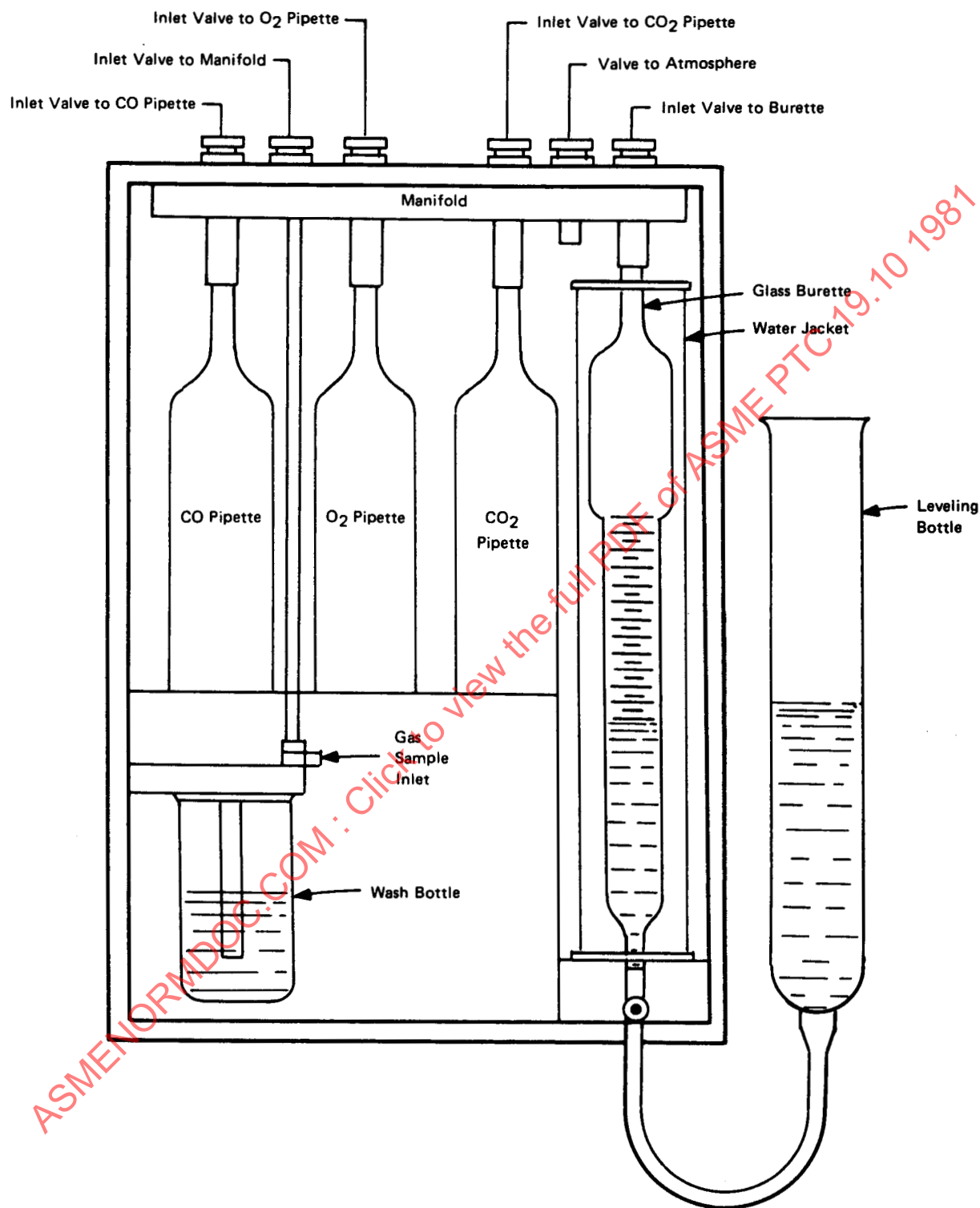


FIG. 13 TYPICAL ORSAT APPARATUS

ASME PERFORMANCE TEST CODES

the pipettes shall be kept free from absorbent solution. One drop is sufficient to spoil the test reading.

6.01.1.3.3 As a safety measure, take care not to let the reagents contact any part of the body or clothing.

6.01.1.3.4 Use water between the water-jacket and burette to prevent variations in gas volume because of changes in sample gas temperature.

6.01.1.3.5 All gases, especially sulfur dioxide and carbon dioxide, are soluble in water to some extent. Bubble the sample rapidly through the confining liquid in the burette for 10 minutes to assure saturation with sample gas before taking readings.

6.01.1.3.6 Absorbent solutions remove constituents chemically and, to a small extent, physically. However, after the first one or two analyses, the solutions become saturated and the error caused by solubility becomes negligible.

6.01.1.3.7 A 5°F (2.8°C) change in temperature of

the gas during analysis makes a difference of 1 percent in volume at constant pressure. This source of error is eliminated by allowing sufficient time for the apparatus to stabilize at ambient temperature before reading and by shielding from drafts.

6.01.1.3.8 Allow time for the solution to drain down the sides of the burette, and this time should be approximately the same before each measurement.

6.01.1.3.9 Leakage tests shall be made before using the apparatus.

6.01.1.3.10 Keep connecting tubing between the sampling probe and Orsat free of leakage and as short as possible.

6.01.1.4 Absorbing Reagents

The chemical reagents normally used in the Orsat apparatus are given in Table 3.

TABLE 3
Standard Orsat Reagents

Component Absorbed	Reagent	Remarks
Carbon Dioxide (CO ₂)	(a) Potassium Hydroxide (b) Sodium Hydroxide	Dissolves 20 times its own volume of CO ₂ .
Oxygen (O ₂)	(a) Alkaline Pyrogalllic Acid (b) Chromous Chloride (c) Fieser's Solution (d) Sodium Pyrogalllic Acid	Dissolves 10 times its own volume of O ₂ . Light sensitive, keep in dark place. Most rapid known. Difficult to prepare. Unstable. Used when sodium hydroxide is used for CO ₂ .
Carbon Monoxide (CO)	(a) Acid Cuprous Chloride (b) Cuprous-Sulphate Betanaphthol	
	Confining Solution	Used as measuring burette and displacement fluid. Minimizes error due to gas solubility.

Standard Test Method for OXIDES OF NITROGEN IN GASEOUS COMBUSTION PRODUCTS (PHENOL-DISULFONIC ACID PROCEDURES)¹

This Standard is issued under the fixed designation D 1608; 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.

1. Scope and Application

1.1 This method describes the phenol-disulfonic acid colorimetric procedure (1)² for the determination of total oxides of nitrogen [nitrous oxide (N₂O) excepted] in gaseous effluents from combustion and other nitrogen oxidation processes.

1.2 It is applicable to a concentration range of oxides of nitrogen as nitrogen dioxide (NO₂) of 5 ppm to several thousand parts per million by volume (four to several thousand milligrams per dry standard cubic metre).

1.3 Since the grab sampling technique used takes a relatively small sample over a very short period of time, the result obtained will be an instantaneous measure of the nitrogen oxides and, therefore, will be representative of the emissions only if the gas stream is well mixed and the concentration constant with time. Multiple samples are recommended.

2. Applicable Documents

2.1 ASTM Standards:

- D 1193 Specification for Reagent Water³
- D 1356 Definitions of Terms Relating to Atmospheric Sampling and Analysis⁴
- D 1357 Recommended Practice for Planning the Sampling of the Atmosphere⁴
- D 1605 Recommended Practices for Sampling Atmospheres for Analysis of Gases and Vapors⁴
- E 200 Preparation, Standardization, and Storage of Standard Solutions for Chemical Analysis⁴

3. Summary of Method

3.1 The gas sample is admitted into an

evacuated flask containing an oxidizing absorbing solution consisting of hydrogen peroxide in dilute sulfuric acid. The oxides of nitrogen are converted to nitric acid by gas phase oxidation due to oxygen in the sample and by the absorbent solution. The resulting nitrate ion is reacted with phenol disulfonic acid to produce a yellow compound (the tri-ammonium salt of 6-nitro-1-phenol-2,4-disulfonic acid) which is measured colorimetrically. Calibration curves, prepared from samples of known nitrate content, are used to determine the amount of nitrate in the sample with results expressed as nitrogen dioxide.

4. Significance

4.1 This method provides a means to measure the total nitrogen oxides (NO_x) content of gaseous emissions for purposes such as determining compliance with regulations, studying the effect of various abatement procedures on NO_x emissions, and checking the validity of instrumental measurements.

5. Definitions

5.1 For definitions of terms used in this method, refer to Definitions D 1356.

6. Interferences (1,2)

6.1 Inorganic nitrates, nitrites, or organic nitrogen compounds that are easily oxidized to nitrates interfere with the method and give erroneously high results. The presence of certain reducing agents, for example, sulfur dioxide (SO₂), may interfere by consuming part of the hydrogen peroxide in the absorbing solution to leave an inadequate amount for reaction with the oxides of nitrogen. Halides lower the results but interference from halide ion (and lead) are negligible in the concentration usually encountered in combustion sources.

6.2 The role of some of the constituents of combustion effluents as possible interfering substances has not been thoroughly investigated.

¹ This method is under the jurisdiction of ASTM Committee D-27 on Sampling and Analysis of Atmospheres and is the responsibility of Subcommittee D22.06 on Source Sampling.

Current edition approved April 29, 1977. Published June 1977. Originally published as D 1608 - 58 T. Last previous edition D 1608 - 60 (1967).

² The boldface numbers in parentheses refer to a list of references at the end of this method.

³ *Annual Book of ASTM Standards*, Part 31.

⁴ 1976 *Annual Book of ASTM Standards*, Part 26.

7. Apparatus

7.1 The assembled sampling apparatus is shown in Fig. 1.

7.2 *Barometer*, capable of measuring atmospheric pressure to ± 2 mm (0.1 in.) Hg.

7.3 *Bottles*, 120-ml or 4-oz, glass or polyethylene, with leak-free noncontaminating caps.

7.4 *Evaporating Dishes*, new condition, unetched borosilicate glass or porcelain, about 200-ml capacity. Do not use platinum ware (7).

7.5 *Mercury Manometer*, open-end U-tube, 1-mm (or 36-in.) with 1-mm (or 0.1-in.) divisions.

NOTE 1—An unbreakable, roll-up type is convenient for this application.

7.6 *Microburet*, 10-ml capacity, with 0.01-ml divisions.

7.7 *pH Paper*, or litmus paper, covering the range from 7 to 14 pH.

7.8 *Sample Collection Flask, Calibrated*—Two-liter round-bottom glass flask with a short neck 24/40 standard-taper joint, protected against implosion or breakage with tape or foamed plastic, with known volume.

7.9 *Sampling Probe*, borosilicate glass, approximately 6-mm inside diameter, fitted with a 12/5 spherical joint for attachment to the three-way stopcock on the sample collection flask, provided with a filter on the inlet end for removal of particulate matter, long enough to extend about third- to halfway into the stack or duct (or at least 1 m beyond inside wall of stacks greater than 2 m in diameter), and heated or insulated, or both, sufficiently well to prevent condensation of moisture while purging and sampling.

7.10 *Spectrophotometer*, or filter photometer, capable of measuring absorbance at 405 nm.

NOTE 2—A wavelength of 400 nm was actually used in the Project Threshold tests, but recent work (3,6,7) has shown that the absorbance peak maximum is actually at 405 nm which is therefore a better choice. This change should tend to improve the precision and accuracy of the method.

7.11 *Squeeze-bulb*, rubber, valved for one-way flow to purge sampling probe.

7.12 *Stirring Rod*, polyethylene, to avoid scratching the evaporating dishes.

7.13 *Stopcock, Three-way, T-bore*, with a 24/40 $\frac{1}{2}$ joint for attachment to the sample collection flask, and a 12/5 spherical joint for attachment to the sampling probe.

7.14 *Stopcock Grease*, high vacuum, high temperature, inert.⁵

⁵ "Halocarbon Chemically Inert Stopcock Grease" made by the Halocarbon Products Corp., 82 Burlews Court, Hackensack, N. J., has been found satisfactory for this purpose.

7.15 *Thermometer*, dial-type or glass, with 1°C (2°F) divisions and an approximate range from -5 to 50°C (25 to 125°F).

7.16 *Vacuum Pump*, portable, capable of evacuating the sample collection flask to a pressure of about 25 mm (1 in.) Hg absolute, or less.

7.17 *Volumetric Flasks*, 50, 100, 1000-ml capacity.

7.18 *Volumetric Pipets*, 1, 2, 3, 4, and 5-ml capacity.

7.19 *Water Bath or Steam Bath*, operating at approximately 100°C (212°F) (an electric hot plate is not suitable because it tends to cause spattering and possible loss of sample).

8. Reagents

8.1 *Purity of Reagents*—Reagent grade chemicals shall be used in all tests. Unless otherwise indicated, it is intended that all reagents shall conform to the specifications of the Committee on Analytical Reagents of the American Chemical Society, where such specifications are available.⁶ Other reagents may be used provided it can be demonstrated that they are of sufficiently high purity to permit their use without decreasing the accuracy of the determination.

8.2 *Purity of Water*—Unless otherwise indicated, references to water shall be understood to mean reagent water conforming to Type III or better of Specification D 1193. Additionally, this method requires water that is nitrate-free as demonstrated by a low blank absorbance reading (less than 0.002 nm) in Section 12.

8.3 *Absorbing Solution*—Dilute 3.0 ml of hydrogen peroxide (H₂O₂, 3 %) to 100 ml with sulfuric acid solution (H₂SO₄, 3+997). A fresh solution must be prepared weekly. Do not expose to excessive heat or direct sunlight for prolonged time.

8.4 *Ammonium Hydroxide (sp gr 0.90)*—Concentrated ammonium hydroxide (NH₄OH). A fresh solution must be used.

8.5 *Fuming Sulfuric Acid (H₂SO₄·XSO₃)*, 15 to 18 weight percent free sulfur trioxide. **Caution**—Handle with care.

8.6 *Hydrogen Peroxide (3 %)*—Dilute 10 ml of concentrated H₂O₂ (30 %) to 100 ml. A fresh solution must be prepared each time new absorbing solution is prepared.

NOTE 3—If the strength of the H₂O₂ (30 %) is in doubt, test as follows: Weigh accurately about 5 ml of the H₂O₂ solution and dilute to exactly 500 ml. To 20 ml of the dilute solution add 20 ml of H₂SO₄.

⁶ "Reagent Chemicals, American Chemical Society Specifications," American Chemical Society, Washington, D. C. For suggestions on testing reagents not listed by ACS, see "Reagent Chemicals and Standards," by Joseph Rosin, D. Van Nostrand Company, Inc., New York, N. Y., and the "United States Pharmacopeia."

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(1+9) and titrate with 0.1 *N* potassium permanganate (KMnO_4) solution to a permanent pink color. One millilitre of 0.1 *N* KMnO_4 solution = 0.001701 g of H_2O_2 .

8.7 *Phenol* ($\text{C}_6\text{H}_5\text{OH}$), pure white solid.

8.8 *Phenol Disulfonic Acid Solution*—Dissolve 25 g of phenol in 150 ml of concentrated H_2SO_4 (sp gr 1.84) by heating on a steam bath. Cool, add 75 ml of fuming H_2SO_4 (15 to 18 % SO_3) and heat on the steam bath at 100°C (212°F) for 2 h. Cool and store in a dark glass-stoppered bottle. The solution should be colorless; it deteriorates on long standing. Discard if a yellow color develops.

8.9 *Potassium Nitrate* (KNO_3)—Dry in an oven at 105 to 110°C for 2 h just before preparation of the standard solution.

8.10 *Potassium Nitrate, Stock Standard Solution* (1 ml = 1 mg NO_2)—Dissolve exactly 2.1980 g of dried KNO_3 in water and dilute to 1 litre in a volumetric flask.

8.11 *Potassium Nitrate, Working Standard Solution* (1 ml = 100 μg NO_2)—Dilute 10 ml of the stock standard potassium nitrate solution (1 ml = 1 mg NO_2) to 100 ml with water in a volumetric flask.

8.12 *Sodium Hydroxide Solution* (~1 *N*)—Slowly add 40 g of sodium hydroxide (NaOH) pellets to 800 to 900 ml of water in a 2-litre beaker with stirring until all pellets are dissolved. Dilute to 1 litre with water and mix well. Store in an airtight polyethylene or polypropylene bottle.

8.13 *Sulfuric Acid* (sp gr 1.84)—Concentrated H_2SO_4 , minimum assay 95 %.

8.14 *Sulfuric Acid* (3+997)—Carefully add 3 ml of concentrated H_2SO_4 (sp gr 1.84) to water and dilute to 1 litre.

9. Safety Precautions

9.1 Cover the glass sample collection flask, which is evacuated during the sampling procedure, with tape or foamed plastic to avoid injury in case of implosion or breakage.

9.2 The fuming sulfuric acid used in preparing the phenol disulfonic acid reagent is highly corrosive and fumes in moist air. Wear protective gloves, apron, and face shield, and carry out the reagent preparation in a hood.

9.3 Concentrated acids and bases are used throughout the laboratory procedure. Use care when adding them to other solutions to avoid overheating and skin contact.

10. Sampling

10.1 Pipet 25.0 ml of absorbing solution into a calibrated sampling flask and attach the three-way stopcock to the flask (Note 4) with the T-bore in the purge position. Insert the sampling probe tip about third- to halfway into the stack or duct, or at least 1 m into

stacks greater than 2 m in diameter, in such a way as to prevent leakage of air into the stack around the probe. Assemble as shown in Fig. 1, making sure that all ground-glass joints have been properly greased, and that all joints and fittings are tight and leak-free. Turn the flask stopcock to the evacuate position and evacuate the flask to the incipient boiling of the solution. If the incipient boiling of the solution cannot be seen, this will be indicated on the manometer by a low and constant reading (usually around 2.5 to 4.0 kPa (20 to 30 torr or 0.8 to 1.2 in. Hg). Shut off the pump valve and then the pump. Check for leakage by observing the manometer for any pressure increase. Any increase greater than 1.33 kPa (10 torr or 0.4 in. Hg) over a 1-min period is not acceptable; do not take a sample until the leakage problem is corrected. Turn the flask stopcock to the purge position. Using a pump or a valved rubber squeeze bulb, thoroughly purge the sampling probe and the flask stopcock with the sample gas until the probe and stopcock are warmed to the gas temperature and free of condensate. If condensation remains, heat the probe and purge until the condensation disappears. With the pump valve and pump shut off, turn the flask stopcock to the evacuate position and record the flask temperature, the difference in mercury levels in the manometer, and the barometric pressure. The absolute pressure in the flask is equal to the barometric pressure minus the manometer reading. Immediately, turn the stopcock to the sample position so that the gas enters the flask and the pressures in the flask and the sample line are equalized (usually about 15 s are sufficient). Turn the stopcock to the purge position to seal the flask and allow the gas to remain in contact with the absorbing solution overnight (Note 5) at room temperature. For further information on sampling, refer to Recommended Practice D 1357 and Recommended Practices D 1605.

NOTE 4—The end 6 mm ($\frac{1}{4}$ in.) of the male 24/40 standard-taper joint is not lubricated, to minimize contact of the gas sample with stopcock grease during absorption.

NOTE 5—If an overnight absorption period is not feasible, the sample can be shaken initially and every 20 min for a 2-h period. The result will not be significantly different than for a static overnight absorption period, provided that the oxygen concentration in the flask is greater than 1 %.

11. Calibration

11.1 *Sample Collection Flask Volume*—Fill the sample flask and stopcock assembly with water up to the stopcock plug. Determine the volume to ± 10 ml by measuring either the volume or weight of the water contained in

the assembly. Number and record the volume on the flask.

11.2 Spectrophotometer Calibration—Prepare a calibration curve to cover a range from about 0 to 125 ppm NO_2 , based on 2000-ml samples of dry gas at 21°C (70°F) and 101.33 kPa (760 torr 29.92 in. Hg). Using a microburet or pipets, transfer 0.0, 1.0, 2.0, 3.0, 4.0, and 5.0 ml of the working standard KNO_3 solution (1 ml = $100\text{ }\mu\text{g NO}_2$) into the 200-ml evaporating dishes and add 25.0 ml of absorbing solution to each. Proceed in accordance with 12.3 to 12.5. Construct a calibration curve by plotting the absorbancies of the solutions at 405 nm against the micrograms of NO_2 .

NOTE 6—Higher sample concentrations may be analyzed using this curve by taking smaller aliquots of the absorbing solution or by dilution, or both, of the color-developed solution before reading in the spectrophotometer.

12. Procedure

12.1 After the absorption period is completed, record the barometric pressure and the room temperature where the sample has stood. Connect one arm of the sample flask stopcock to the open-end manometer, turn the stopcock to open the flask to the manometer, and read the difference between the mercury levels in the manometer. The absolute internal pressure in the flask is then the barometric pressure less this difference. Correct the gas volume in the flask as directed in 13.1.

12.2 Transfer the absorbing solution quantitatively from the flask into a 200-ml evaporating dish (Note 7). Pipet 25.0 ml of unused absorbing solution into another evaporating dish for a blank and add the same amount of water to this dish as was used in transferring the sample. Proceed with the blank in the same manner as directed for the sample.

NOTE 7—If the sample is expected to have a high concentration of oxides of nitrogen, transfer it to a 50-ml volumetric flask instead of the evaporating dish, and dilute to the mark with water. Select a suitable aliquot and pipet it into a 200-ml evaporating dish. Likewise dilute 25.0 ml of unused absorbing solution to 50 ml and pipet an aliquot equal to that of the sample into a 200-ml evaporating dish for a blank.

NOTE 8—To save time a 15-ml aliquot of the used absorbing solution may be pipetted into the evaporating dish, rather than quantitatively transferring the whole solution. This is permissible if the concentration of nitrogen dioxide (NO_2) is greater than 10 ppm, and if the evacuation of the flask prior to admitting the sample was to the incipient or flash boiling point of the unused absorbing solution. Evacuation up to 1 min after flash boiling appears to result in about 1 % decrease in the volume of the solution.

NOTE 9—If necessary, in order to immediately reuse the sampling flask or to ship the used absorbing solution back to a laboratory for analysis, the entire used absorbing solution or a 15-ml aliquot may be quantitatively transferred to a leak-free glass or polyethylene bottle until ready for analysis.

12.3 Add NaOH solution to the sample solution to the evaporating dish and to the blank until each is just basic to litmus or pH paper. Do not add any excess NaOH (7). Evaporate each to dryness on the water bath and allow to cool. Very carefully add 2 ml of phenol disulfonic acid solution to each residue and triturate thoroughly with a polyethylene rod to ensure complete contact of the residue with the solution. Add 1 ml of water and 4 drops of H_2SO_4 (sp gr 1.84) to each and heat on the water bath for 3 min with occasional stirring. Allow the mixture to cool, add 10 ml of water to each, and mix well by stirring. Add 15 ml of fresh, cool NH_4OH dropwise to each, with constant stirring. Test with litmus paper to make sure an excess of the NH_4OH is present.

12.4 Filter the solutions through 7-cm, rapid, medium-texture filter papers (Note 10) into 50-ml volumetric flasks. Wash the evaporating dishes three times with 4 to 5 ml of water and pass the washing through the filters. Make up the volumes of the solutions to 50 ml with water and mix thoroughly.

NOTE 10—Reference (7) recommends discarding any sample for which a significant precipitate is present or the evaporating dish is visually etched. However, the need for this has not been demonstrated. The same grade filter paper should, however, be used both in preparing the calibration curves and in running the samples. It has been found that some yellow color is retained on the paper when filtering more concentrated samples and this factor must be taken into account by use of the sample type of filter paper throughout, or by continued washing until no color is retained in any case. Alternatively, the ammoniacal solutions may be centrifuged, instead of filtered, after dilution to 50 ml.

12.5 Read the absorbance of the sample solution against the blank in suitable equipment for measurement at 405 nm. If the absorbance falls beyond the range of calibration, thinner cells may be used or a suitable aliquot selected. Dilute the aliquot and the blank to the same volume and read the absorbance of the sample aliquot against that of the blank aliquot.

12.6 Convert the photometric readings to micrograms of NO_2 by means of the calibration curves.

NOTE 10—The calibration curves must not be assumed to be usable over any protracted length of time. One or more standards should be run along with the samples each time a set is run, or at least every few days if samples are being run daily.

13. Calculations

13.1 Gas Sample Volume Correction—Correct the volume of the dry gas sample (after removal of absorbed constituents) to 21°C (70°F) and 101.33 kPa (760 torr or 29.92 in. Hg).

13.1.1 Calculate the volume of gas sample as follows:

$$V_c = (V_t - V_a)(P_i/T_i - P_f/T_f)(21 + 273.2)/101.33$$

where:

V_c = corrected volume of sample gas, ml,

V_t = volume of sample collection flask, ml,

V_a = volume of absorbing solution used, 25 ml,

P_i = initial absolute pressure in flask prior to sampling, kPa,

P_f = final absolute pressure in flask after absorption period, kPa,

T_i = initial absolute temperature of flask prior to sampling, K ($^{\circ}\text{C} + 273.2$), and,

T_f = final absolute temperature of flask after absorption period, K ($^{\circ}\text{C} + 273.2$).

13.2 Calculate the concentration of NO_2 in parts per million by volume (ppm), dry basis, as follows:

$$\text{NO}_2, \text{ ppm} = (24.1 W \times 10^3)/46V_c$$

where:

V_c = corrected volume of sample gas, ml,

W = weight of NO_2 found in gas sample μg ,

24.1×10^3 = standard molar gas volume at 21°C (70°F) and 101.33 kPa (760 torr or 29.92 in. Hg), ml, and

46 = formula weight of NO_2 .

NOTE 10—Parts per million by volume of NO_2 may be converted to milligrams per cubic metre (mg/m^3) or to pounds per cubic foot (lb/ft^3) by multiplying by 1.91 and by 1.19×10^{-3} , respectively.

14. Precision and Accuracy (3,4,5)

14.1 In an ASTM-sponsored collaborative effort, laboratories, who had not worked under such conditions before, performed a series of pilot-plant tests indicating a correlation between the precision and the square root of the mean concentration of the NO_x over the range from 20 to 2000 ppm. The measures of precision used presented standard deviations for the components of variance between-laboratories (S_R) and within-laboratories (S_W) obtained from the analysis of variance technique. These correlations are shown in Fig. 2 (5).

14.2 These same laboratories, working in the field at an oil-fired power plant and cement kiln showed a mean between-laboratory standard error of 14.5 ppm over NO_x concentrations in the range from 90 to 260 ppm. The mean between-laboratory standard error obtained from the field data is significantly lower than the comparable standard error measure calculated from the pilot plant work. Presumably, the increased experience of the opera-

tors contributed to the improved precision.

14.3 In order to use these data to determine the 95 % confidence limits (95 % CL) of the mean of n replicate analyses by a single laboratory, the mean level, m , is calculated and the appropriate value of S_W is found from Fig. 2. The 95 % CL of the mean is $m \pm (1.95/\sqrt{n}) \times S_W$.

14.4 The 95 % CL of the estimate that the mean of n replicate analysis by one laboratory represents the mean of a similar number of analyses by a large number of laboratories is calculated from the standard error, S_T , which in turn, can be calculated from the data in Fig. 2 using the equation:

$$S_T = \sqrt{(S_R^2 + S_W^2)/n}$$

If the experience of the laboratories with the method is appropriate and the NO_2 levels is used:

$$S_T = 14.5 \text{ ppm NO}_2$$

Finally, the 95% CL of a single laboratory's data with respect to a mean value determined by many laboratories is $\pm 1.96 \times S_T$.

14.5 In the same collaborative study it was shown that the method had no significant bias over the range studied. Thus, it can be stated that the accuracy of a single laboratory's analysis is $\pm 1.96 \times S_T$ from 20 to 2000 ppm of NO_2 .

14.6 Analysis of standard nitrate solutions following each test indicated that variations introduced by the color formation and measurement portion of the method accounts for roughly 2/3 of the total observed variability. In addition, predominantly negative biases were associated with analysis of these standards ranging from -3.6 to -7.7 % of the true value. Although not statistically significant, the pilot plant spiking tests also produced a negative bias of -3.25 %, suggesting that there was an influence of a negative bias in the chemical portion of the method on the overall results.

14.7 Additional collaborative data concerning precision and accuracy of the phenol-disulfonic acid method, as applied in analysis of effluents by the EPA Method 7, are given in Refs (3) and (4). Generally, these studies showed significantly better precision than indicated by Fig. 2 at NO_x concentrations lower than 400 ppm. The difference in findings may well represent differences in experience of the collaborating laboratories since the field data of the ASTM teams, which were obtained after the initial pilot plant work, was within the range of significance of the EPA correlations.

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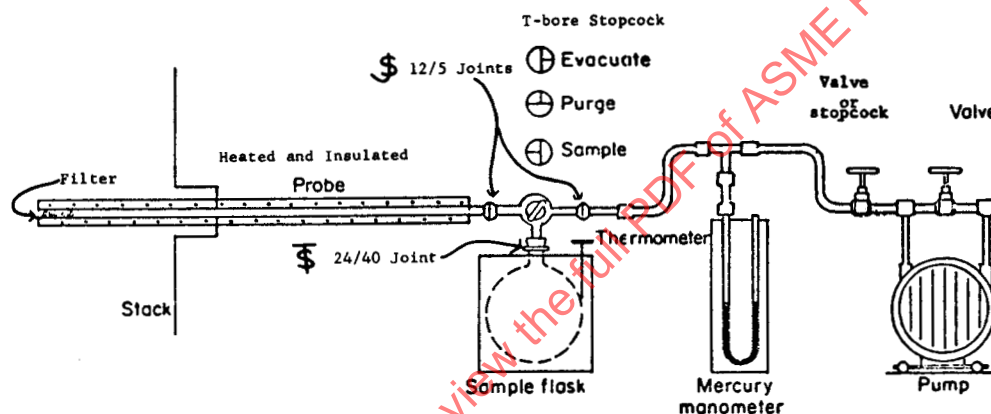


FIG 1 Diagram of Typical Sampling Apparatus used for Determination of Oxides of Nitrogen by ASTM D 1608.

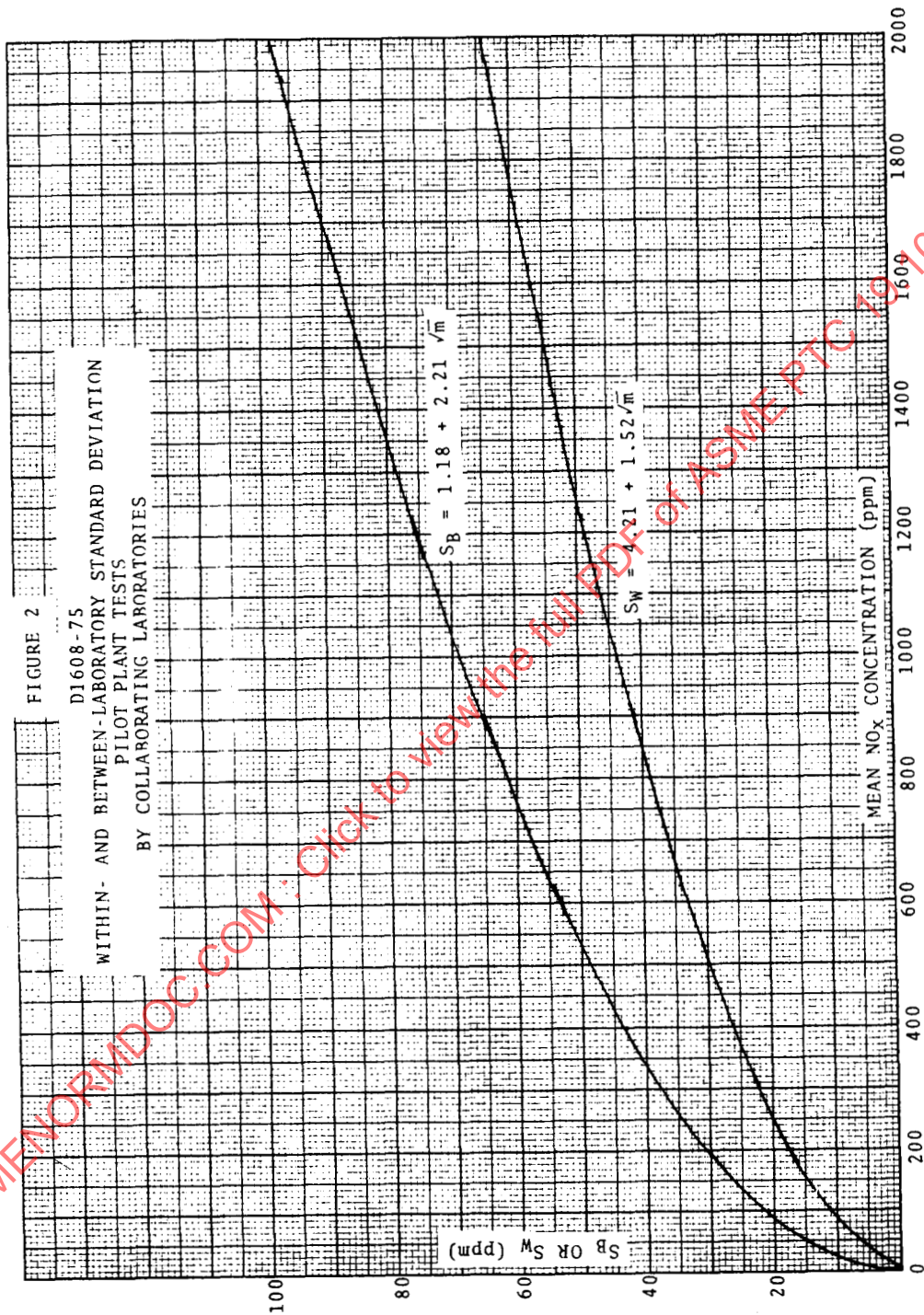


FIG 2 Within- and Between-Laboratory Standard Deviation Pilot Plant Tests by Collaborating Laboratories.

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6.01.3

NITROGEN DIOXIDE IN AIR

Physical and Chemical Analysis Branch

Analytical Method

Analyte:	Nitrogen Dioxide	Method No.:	P&CAM 108
Matrix:	Air	Range:	.01 to 10 $\mu\text{g/liter}$
Procedure:	Colorimetric	Precision:	< 5% RSD
Date Issued:	12/11/72		
Date Revised:	January 10, 1974	Classification:	C (Tentative)

1. Principle of the Method

The nitrogen dioxide is absorbed in an azo dye forming reagent (Reference 1). A stable pink color is produced within 15 minutes which may be read visually or in an appropriate instrument at 550 nm.

2. Range and Sensitivity

- 2.1 This method is intended for the manual determination of nitrogen dioxide in the atmosphere in the range of 0.005 to about 5 parts per million (ppm) by volume or 0.01 to 10 $\mu\text{g/liter}$, when sampling is conducted in fritted bubblers. The method is preferred when high sensitivity is needed.
- 2.2 Concentrations of 5 to 100 ppm in industrial atmospheres and in gas burner stacks also may be sampled by employing evacuated bottles or glass syringes. For higher concentrations, for automotive exhaust, and/or for samples relatively high in sulfur dioxide content, other methods should be applied.*

3. Interferences

- 3.1 A tenfold ratio of sulfur dioxide to nitrogen dioxide produces no effect. A 30-fold ratio slowly bleaches the color to a slight extent. The addition of 1 percent acetone to the reagent before use retards the fading by forming another temporary product with sulfur dioxide. This permits reading within 4 to 5 hours (instead of the 45 minutes required when acetone is not added) without appreciable interferences. Interference from sulfur dioxide may be a problem in some stack gas samples (Section 2.2).

* NOTE: To measure total NO and NO₂. An oxidizing tube must be used to oxidize any NO to NO₂ prior to analysis. At least 3 passes through KGO₃ solution is required.

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- 3.2 A fivefold ratio of ozone to nitrogen dioxide will cause a small interference, the maximal effect occurring in 3 hours. The reagent assumes a slightly orange tint.
- 3.3 Peroxyacynitrate (PAN) can give a response of approximately 15 to 35 percent of an equivalent molar concentration of nitrogen dioxide (Reference 2). In ordinary ambient air the concentrations of PAN are too low to cause any significant error.
- 3.4 The interferences from other nitrogen oxides and other gases which might be found in polluted air are negligible. However, if the evacuated bottle or syringe method is used to sample concentrations above 5 ppm, interference from NO (due to oxidation to NO₂) is possible (Section 8.2.3).
- 3.5 If strong oxidizing or reducing agents are present, the colors should be determined within 1 hour, if possible, to minimize any loss.

4. Precision and Accuracy

- 4.1 A precision of 1 percent of the mean can be achieved with careful work (Reference 3); the limiting factors are the measurements of the volume of the sample and of the absorbance of the color.
- 4.2 At present, accuracy data are not available.

5. Advantages and Disadvantages of the Method

This is a simple method with direct coloration of absorbing reagent, which can be put directly into cuvettes and read, or diluted with absorbing reagent and read.

6. Apparatus

- 6.1 **Absorber.** The sample is absorbed in an all-glass bubbler with a 60 μ maximum pore diameter frit.
 - 6.1.1 The porosity of the fritted bubbler, as well as the sampling flow rate, affect absorption efficiency. An efficiency of over 95 percent may be expected with a flow rate of 0.4 liter/minute or less and a maximum pore diameter of 60 μ . Frits having a maximum pore diameter less than 60 μ will have a higher efficiency but will require an inconvenient pressure drop for sampling (see formula in Section 6.1.2). Considerably lower efficiencies are obtained with coarser frits, but these may be utilized if the flow rate is reduced.

- 6.1.2 Since the quality control by some manufacturers is rather poor, it is desirable to measure periodically the porosity of an absorber as follows: Carefully clean the apparatus with dichromate-concentrated sulfuric acid solution and then rinse it thoroughly with distilled water. Assemble the bubbler, add sufficient distilled water to barely cover the fritted portion, and measure the vacuum required to draw the first perceptible stream of air bubbles through the frit. Then calculate the maximum pore diameter as follows:

$$\text{Maximum pore diameter } (\mu) = \frac{30 s}{P}$$

where:

s = surface tension of water at the test temperature in dynes/cm
(73 at 18°C, 72 at 25°C, and 71 at 31°C),

P = measured vacuum, mm Hg.

- 6.1.3 Rinse the bubbler thoroughly with water and allow to dry before using. A rinsed and reproducibly drained bubbler may be used if the volume (r) of retained water is added to that of the absorbing reagent for the calculation of results. This correction may be determined as follows: Pipet into a drained bubbler exactly 10 ml of a colored solution (such as previously exposed absorbing reagent) of absorbance A_1 . Assemble the bubbler and rotate to rinse the inside with the solution. Rinse the fritted portion by pumping gently with a rubber bulb. Read the new absorbance, A_2 , of the solution. Then:

$$10A_1 = (10 + r) A_2$$

$$\text{or: } r = 10 \frac{A_1}{A_2} - 1$$

- 6.2 **Air Metering Device.** A glass rotameter capable of accurately measuring a flow of 0.4 liter/minute is suitable. A wet test meter is convenient to check the calibration.

- 6.3 **Sampling Probe.** A glass or stainless steel tube 6 to 10 mm in diameter provided with a downward-facing intake (funnel or tip) is suitable. A small loosely fitting plug of glass wool may be inserted, when desirable, in the probe to exclude water droplets and particulate matter. The dead volume of the system should be kept minimal to permit rapid flushing during sampling to avoid losses of nitrogen dioxide on the surfaces.

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- 6.4 **Grab-Sample Bottles.** Ordinary glass-stoppered borosilicate glass bottles of 30 to 250 mL sizes are suitable if provided with a mating ground joint attached to a stopcock for evacuation. Calibrate the volume by weighing with connecting piece, first empty, then filled to the stopcock with distilled water.
- 6.5 **Glass Syringes.** 50 or 100 mL syringes are convenient (although less accurate than bottles) for sampling.
- 6.6 **Air Pump.** A vacuum pump capable of drawing the required sample flow for intervals of up to 30 minutes is suitable. A tee connection at the intake is desirable. The inlet connected to the sampling train should have an appropriate trap and needle valve, preferably of stainless steel. The second inlet should have a valve for bleeding in a large excess flow of clean air to prevent condensation of acetic acid vapors from the absorbing reagent, with consequent corrosion of the pump. Alternatively, soda lime may be used in the trap. A filter and critical orifice may be substituted for the needle valve (Reference 4).
- 6.7 **Spectrophotometer or Colorimeter.** A laboratory instrument suitable for measuring the pink color at 550 nm, with stoppered tubes or cuvettes. The wavelength band width is not critical for this determination.
- 6.8 Assorted laboratory glassware.
7. **Reagents**
- All chemicals used should be analytical reagent grade (Reference 5).
- 7.1 **Nitrite-Free Water.** All solutions are made in nitrite-free water. If available distilled or deionized water contains nitrite impurities (produces a pink color when added to absorbing reagent), redistill it in an all-glass still after adding a crystal each of potassium permanganate and of barium hydroxide.
- 7.2 **N-(1-Naphthyl) -Ethylenediamine Dihydrochloride, Stock Solution (0.1%).** Dissolve 0.1 g of the reagent in 100 mL of water. Solution is stable for several months if kept well stoppered in a brown bottle in the refrigerator. (Alternatively, weighed small amounts of the solid reagent may be stored.)
- 7.3 **Absorbing Reagent.** Dissolve 5 g of anhydrous sulfanilic acid (or 5.5 g of $\text{NH}_2\text{C}_6\text{H}_4\text{SO}_3\text{H}\cdot\text{H}_2\text{O}$) in almost a liter of water containing 140 mL of glacial acetic acid. Gentle heating is permissible to speed up the process. To the cooled mixture, add 20 mL of the 0.1% stock solution of N-(1-naphthyl) -ethylenediamine dihydrochloride, and dilute to 1 liter. Avoid lengthy contact with air during both preparation and use, since discoloration of reagent will result because of absorption of nitrogen dioxide. The solution is stable for several months if kept

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well-stoppered in a brown bottle in the refrigerator. The absorbing reagent should be allowed to warm to room temperature before use.

- 7.4 Standard Sodium Nitrite Solution (0.0203 g/liter).** One ml of this working solution of sodium nitrite (NaNO_2) produces a color equivalent to that of $10\ \mu\text{l}$ of nitrogen dioxide (10 ppm in 1 liter of air at 760 mm of mercury and 25°C : see Section 10.2.1). Prepare fresh just before use by dilution from a stronger stock solution containing 2.03 g of the reagent grade granular solid (calculated as 100 percent) per liter. It is desirable to assay (Reference 5), the solid reagent, especially if it is old. The stock solution is stable for 90 days at room temperature, and for a year in a brown bottle under refrigeration.

8. Procedure

- 8.1 Cleaning of Glassware.** All washed glassware should be allowed to stand awhile in chromic acid solution, and then thoroughly rinsed with single and then double distilled water.
- 8.2 Collection and Storage of Samples.** Three methods are described below. Concentrations below 5 ppm are sampled by the bubbler method. Higher concentrations may be sampled by the evacuated bottle method, or more conveniently (but less accurately) by the glass syringe method. The latter method is more useful when appreciable concentrations (e.g., 20 ppm) of nitric oxide are expected.

8.2.1 Bubbler Method

1. Assemble, in order, a sampling probe (optional), a glass rotameter, fritted absorber, and pump. Use ground-glass connections upstream from the absorber. Butt-to-butt glass connections with slightly greased vinyl or pure gum rubber tubing also may be used for connections without losses if lengths are kept minimal. The sampling rotameter may be used upstream from the bubbler provided occasional checks are made to show that no nitrogen dioxide is lost. The rotameter must be kept free from spray or dust.
2. Pipet 10.0 ml of absorbing reagent into a dry fritted bubbler (Section 6.1.3).
3. Draw an air sample through it at the rate of 0.4 liter/minute (or less) long enough to develop sufficient final color (about 10 to 30 minutes). Note the total volume of air sampled.
4. Measure and record the sample air temperature and pressure.

8.2.2 Evacuated Bottle Method

1. Sample in bottles of appropriate size containing 10.0 mL (or other convenient volume) of absorbing reagent. For 1 cm spectrophotometer cells, a 5:1 ratio of air sample volume to reagent volume will cover a concentration range up to 100 ppm; a 25:1 ratio suffices to measure down to 2 ppm.
2. Wrap a wire screen or glass-fiber-reinforced tape around the bottle for safety purposes.
3. Grease the joint lightly with silicone or fluorocarbon grease.
4. If a source of vacuum is available at the place of sampling, it is best to evacuate just before sampling to eliminate any uncertainty about loss of vacuum. A three-way Y stopcock connection is convenient. Connect one leg to the sample source, one to the vacuum pump, and the third to a tee attached to the bottle and to a mercury manometer or accurate gauge. In the first position of the Y stopcock, the bottle is evacuated to the vapor pressure of the absorbing reagent. In the second position of the Y stopcock the vacuum pump draws air through the sampling line to flush it thoroughly. The actual vacuum in the sample bottle is read on the manometer. In the third position of the Y stopcock the sampling line is connected to the evacuated bottle and the sample is collected.
5. The stopcock on the bottle is then closed. Allow 15 minutes with occasional shaking for complete absorption and color development.
6. For calculation of the standard volume of the sample, record the temperature and the pressure. The pressure is the difference between the filled and evacuated conditions. The uncorrected volume is that of the bottle plus that of the connection up to the stopcock minus the volume of absorbing reagent.

8.2.3 Glass Syringe Method

1. Ten mL of absorbing reagent is kept in a capped 50 (or 100) mL glass syringe, and 40 (or 90) mL of air is drawn in at the time of sampling.
2. The absorption of nitrogen dioxide is completed by capping and shaking vigorously for 1 minute, after which the air is expelled. (When appreciable concentrations -- e.g., 20 ppm -- of nitric oxide are suspected, interference caused by the oxidation of nitric oxide to nitrogen dioxide is minimized by expelling the air sample immediately after the absorption period.)

3. Additional air may be drawn in and the process repeated several times, if necessary, to develop sufficient final color.

8.2.4 Effects of Storage. Colors may be preserved, if well stoppered, with only 3 to 4% loss in absorbance per day; however, if strong oxidizing or reducing gases are present in the sample in concentrations considerably exceeding that of the nitrogen dioxide, the colors should be determined as soon as possible to minimize any loss. (See Section 3 for effects of interfering gases.)

8.3 Analysis of Samples

8.3.1 After collection or absorption of the sample, a red-violet color appears. Color development is complete within 15 minutes at room temperature.

8.3.2 Compare with standards visually or transfer to stoppered cuvettes and read in a spectrophotometer at 550 mμ, using unexposed reagent as a reference. Alternatively, distilled water may be used as a reference and the absorbance of the reagent blank deducted from that of the sample.

8.3.3 Colors too dark to read may be quantitatively diluted with unexposed absorbing reagent. The measured absorbance is then multiplied by the dilution factor.

9. Calibration and Standardization

Either of two methods of calibration may be employed. The most convenient is standardizing with nitrite solution. Greater accuracy is achieved by standardizing with accurately known gas samples in a precision flow dilution system (References 3,6,7). The recently developed permeation tube technique (Reference 8) appears promising. If the gaseous method is used, the stoichiometric factor is eliminated from the calculations. Concentrations of the standards should cover the expected range of sample concentrations.

9.1 Nitrite Solution Method

9.1.1 Add graduated amounts of the NaNO₂ solution up to 1 ml (measured accurately in a graduated pipet or small buret) to a series of 25 ml volumetric flasks, and dilute to the mark with absorbing reagent.

9.1.2 Mix, allow 15 minutes for complete color development and read the colors (Section 8.3).

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9.1.3 Good results can be obtained with these small volumes of standard solution if they are carefully measured. Making the calibration solutions up to 25 ml total volume, rather than the 10 ml volume used for samples, facilitates accuracy. If preferred, even larger volumes may be used with correspondingly larger volumetric flasks.

9.1.4 Plot the absorbances of the standard colors against the μl of nitrogen dioxide per ml of absorbing reagent. The latter values are equal to the corresponding ml of standard nitrite solution times 0.4 (Section 10.2.2). If preferred, transmittance may be plotted instead of absorbance, using semilogarithmic graph paper. The plot follows Beer's law. Draw the straight line through the origin giving the best fit, and determine the slope, K, which is μl of NO_2 intercepted at absorbance of exactly 1.0 or at 10 percent transmittance. For 1 cm cells, the value of K is about 0.73.

9.2 **Gaseous Standard Methods.** Two techniques are outlined below. Consult the original references for complete details, and Section 10.3 for calculations.

9.2.1 Method 1 (Reference 3) for gaseous standardization is as follows:

1. About 5 ml of pure liquid nitrogen dioxide is placed in a small glass bubbler (10 mm in diameter and 100 mm in length) provided with ground glass stopcocks and spherical joints on both intake and outlet tubes.
2. The bubbler is immersed in a thermos bottle ice bath and connected to an air line. A small pump with flowmeter delivers a steady stream of a few ml/min of air to the bubbler, thence through two flowmeters which permit the discarding of up to 90 percent of the NO_2 and finally to a large stream of carbon filtered air (1000 to 1500 liters/min) from a small blower. All this air passes through a 10 cm Biram anemometer mounted on the end of a pipe 10 cm in diameter. It has been found empirically with this arrangement that the anemometer reading in ft/min times 1.64 is equivalent to liters/min.
3. The bubbler is weighed to 0.1 mg at the beginning and end of an accurately measured time period. The stopcocks are closed each time before the bubbler is removed from the ice bath and wiped dry for weighing.

9.2.2 Method 2 (References 6,7) for preparing known dilutions of nitrogen dioxide consists in making a preliminary dilution (about 0.4%) of nitrogen dioxide in air in a stainless steel tank at 1000 lbs pressure. Subsequent dilution, by air in a flow system at atmospheric pressure, of the analyzed

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tank mixture controlled by an asbestos plug and manometer yields concentrations of 0.1 to 10 ppm NO₂

- 9.2.3 Sample the gas mixtures by the bubbler method (Section 8.2.1), and read the colors (Section 8.3). Select concentrations and sample volumes to produce colors covering the accurate absorbance range of the spectrophotometer.
- 9.2.4 Standardization by gaseous samples can be based either upon a weight-volume relationship if the source of nitrogen dioxide is weighed, or a volume-volume relationship if the source is an analyzed tank mixture.
- 9.2.5 Calculate the concentration of the sample air stream, C , in ppm by volume ($\mu\text{l NO}_2/\text{l of air}$).

$$C = \frac{10^6 (W_1 - W_2)}{t} \times \frac{0.532}{F_1} \times \frac{F_2}{F_3}$$

or: $C = C_t \times \frac{F_2}{F_3}$

where:

- C = concentration of sample air stream in ppm.
- $(W_1 - W_2)$ = difference between initial and final weights, in grams, of nitrogen dioxide bubbler (or of permeation tube); weight loss is usually 0.01 to 0.05 g.
- t = time interval in minutes, between the weighings.
- 0.532 = (24.47/46.0), ideal volume in l at 25°C and 760 mm Hg of 1.0 g of nitrogen dioxide.
- F_1 = flow rate, in lpm, of air passed through nitrogen dioxide bubbler (or over permeation tube), corrected to 25°C and 760 mm Hg.
- F_2 = flow rate, in lpm, of concentrated gas mixture injected into sample air stream.
- F_3 = total flow rate, in lpm, of sample air stream
- C_t = analyzed concentration of the tank mixture, in ppm by volume on an ideal gas basis.

- 9.2.6 For each standard color, calculate the μl of nitrogen dioxide/ml of absorbing reagent:

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$$\mu\text{l NO}_2/\text{ml} = C V$$

where:

C = ppm concentration (Section 9.2.5)

V = volume of air sample at 25°C and 760 mm Hg, in l/ml of absorbing reagent.

9.2.7 Plot the absorbances of the colors against the μl of gaseous nitrogen dioxide/ ml of absorbing reagent. Draw the straight line through the origin giving the best fit, and determine the slope, K (the value of $\mu\text{l/ml}$ intercepted at absorbance of exactly 1.0).

10. Calculations

10.1 For convenience, standard conditions are taken as 760 mm Hg and 25°C, at which the molar gas volume is 24.47 liters. (This is identical with standard conditions for Threshold Limit Values of the American Conference of Governmental Industrial Hygienists; it is very close to the standard conditions used (Reference 9) for air-handling equipment, of 29.92 in. Hg, 70°F, and 50 percent relative humidity, at which the molar gas volume is 24.76 liters, of 1.2 percent greater.)

Ordinarily the correction of the sample volume to these standard conditions is slight and may be omitted; however, for greater accuracy, it may be made by means of the perfect gas equation.

10.2 Standardization by nitrite solution is based upon the empirical observation (References 1,6) that 0.72 mole of sodium nitrite produces the same color as 1 mole of nitrogen dioxide.

10.2.1 This factor is applied to calculate the equivalence of the nitrite solution to the volume of NO_2 absorbed as follows: One ml of the working standard solution contains 2.03×10^{-5} g NaNO_2 . Since the molecular weight of NaNO_2 is 69.00, this is equivalent to:

$$\frac{2.03 \times 10^{-5}}{69.00} \times \frac{24.47}{0.72} = 1 \times 10^{-5} \text{ l of NO}_2, \text{ or}$$
$$= 10 \mu\text{l of NO}_2$$

10.2.2 In Section 9.1, the calibration standard containing 1 ml of nitrite solution ($10 \mu\text{l NO}_2$) per 25 ml total volume is equivalent to 10/25 or 0.4 μl of NO_2 per ml .

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10.3 Compute the concentration of nitrogen dioxide in the sample as follows:

$$\text{Nitrogen dioxide, ppm} = AK/V$$

where:

A = measured absorbance

K = standardization factor from Section 9.1.4 or 9.2.7

V = volume of air sample, at 25°C and 760 mm Hg, in liters/ml of absorbing reagent.

10.4 If preferred, the graph from Section 9.1.4 or 9.2.7 may be used instead as follows:

$$\text{Nitrogen dioxide, ppm} = \mu\ell \text{ NO}_2 \text{ per ml/V}$$

10.4.1 If V is a simple multiple of K, calculations are simplified. Thus, for the K value of 0.73 previously cited, if exactly 7.3 liters of air are sampled through a bubbler containing 10 ml of absorbing reagent, K/V = 1, and the absorbance is also ppm directly.

10.4.2 For exact work, an allowance may be made in the calculations for sampling efficiency and for fading of the color using the following equation:

$$\text{Nitrogen dioxide, ppm} = A_c K/VE$$

where:

A_c = corrected absorbance; the absorbance is corrected for fading of the color as indicated in Section 8.2.4 when there is a prolonged interval between sampling and measurement of the absorbance.

E = sampling efficiency; for a bubbler, E is estimated from prior tests using two absorbers in series (Reference 7 and Section 6.1.1); for a bottle or syringe, E = 1.0.

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6.01.4

FEDERAL EPA METHOD

Determination of Sulfur Dioxide Emissions from
Stationary Sources

1. Principle and Applicability

1.1 Principle. A gas sample is extracted from the sampling point in the stack. The sulfuric acid mist (including sulfur trioxide) and the sulfur dioxide are separated. The sulfur dioxide fraction is measured by the barium-thorin titration method.

1.2 Applicability. This method is applicable for the determination of sulfur dioxide emissions from stationary sources. The minimum detectable limit of the method has been determined to be 3.4 milligrams (mg) of SO_2/m^3 (2.12×10^{-7} lb/ft³). Although no upper limit has been established, tests have shown that concentrations as high as 80,000 mg/m³ of SO_2 can be collected efficiently in two midjet impingers, each containing 15 milliliters of 3 percent hydrogen peroxide, at a rate of 1.0 lpm for 20 minutes. Based on theoretical calculations, the upper concentration limit in a 20-liter sample is about 93,300 mg/m³.

Possible interferents are free ammonia, water-soluble cations, and fluorides. The cations and fluorides are removed by glass wool filters and an isopropanol bubbler, and hence do not affect the SO_2 analysis. When samples are being taken from a gas stream with high concentrations of very fine metallic fumes (such as inlets to control devices), a high-efficiency glass fiber filter must be used in place of the glass wool plug (i.e., the one in the probe) to remove the cation interferents.

Free ammonia interferes by reacting with SO_2 to form particulate sulfite and by reacting with the indicator. If free ammonia is present (this can be determined by knowledge of the process and noticing white particulate matter in the probe and isopropanol bubbler), alternative methods, subject to the approval of the Administrator, U.S. Environmental Protection Agency, are required.

2. Apparatus

2.1 Sampling. The sampling train is shown in Fig. 6-1, and component parts are discussed below. The tester has the option of substituting sampling equipment described in Method 8 in place of the midjet impinger equipment of Method 6. However, the Method 8 train must be modified to include a heated filter between the probe and

isopropanol impinger, and the operation of the sampling train and sample analysis must be at the flow rates and solution volumes defined in Method 8.

The tester also has the option of determining SO_2 simultaneously with particulate matter and moisture determinations by (1) replacing the water in a Method 5 impinger system with 3 percent peroxide solution, or (2) by replacing the Method 5 water impinger system with a Method 8 isopropanol-filter-peroxide system. The analysis for SO_2 must be consistent with the procedure in Method 8.

2.1.1 Probe. Borosilicate glass, or stainless steel (other materials of construction may be used, subject to the approval of the Administrator), approximately 6-mm inside diameter, with a heating system to prevent water condensation and a filter (either in-stack or heated out-stack) to remove particulate matter, including sulfuric acid mist. A plug of glass wool is a satisfactory filter.

2.1.2 Bubbler and Impingers. One midjet bubbler, with medium-coarse glass frit and borosilicate or quartz glass wool packed in top (see Fig. 6-1) to prevent sulfuric acid mist carryover, and three 30-ml midjet impingers. The bubbler and midjet impingers must be connected in series with leak-free glass connectors. Silicone grease may be used, if necessary, to prevent leakage.

At the option of the tester, a midjet impinger may be used in place of the midjet bubbler.

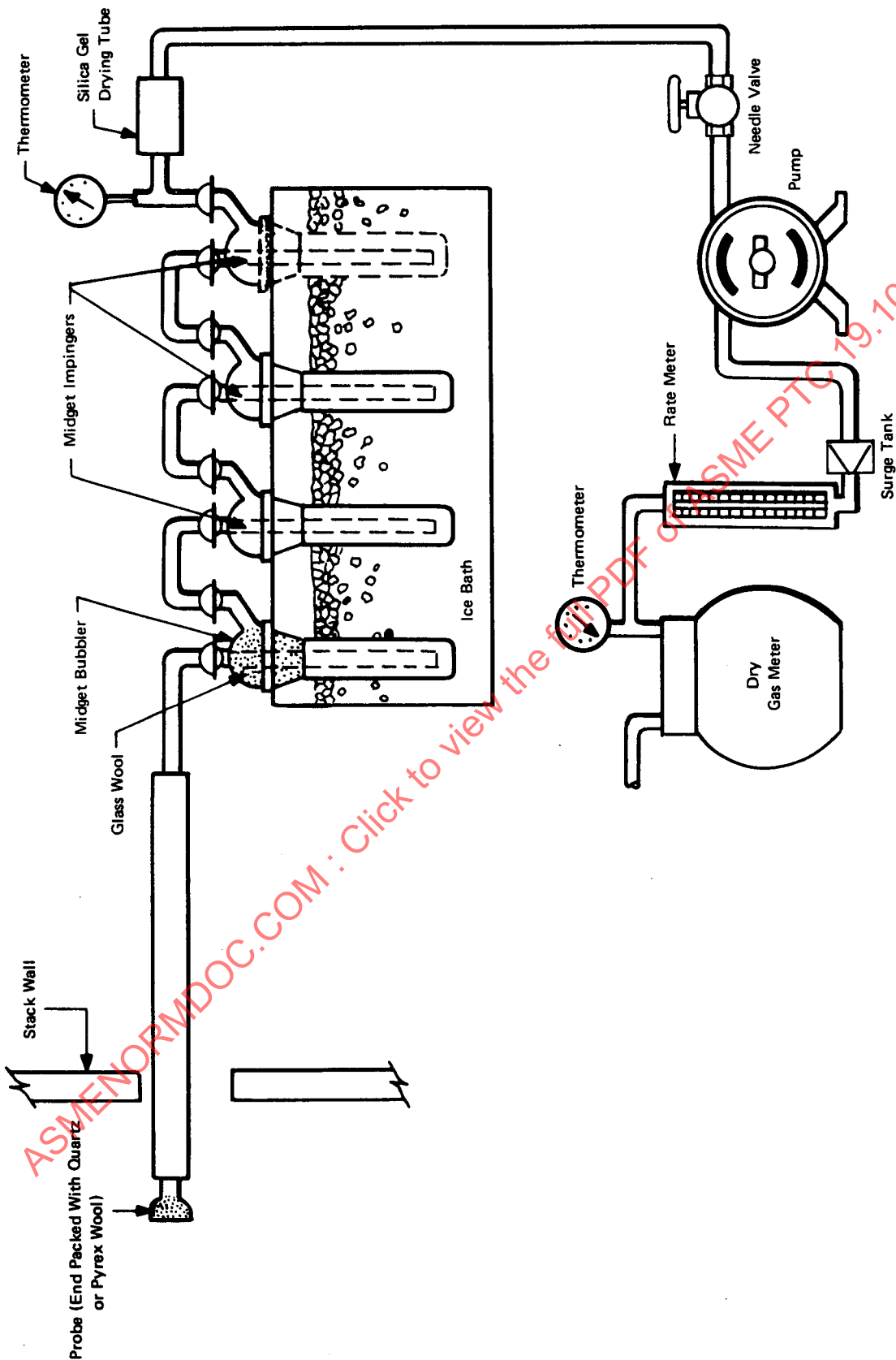
Other collection absorbers and flow rates may be used, but are subject to the approval of the Administrator. Also, collection efficiency must be shown to be at least 99 percent for each test run and must be documented in the report. If the efficiency is found to be acceptable after a series of three tests, further documentation is not required. To conduct the efficiency test, an extra absorber must be added and analyzed separately. This extra absorber must not contain more than 1 percent of the total SO_2 .

2.1.3 Glass Wool. Borosilicate or quartz.

2.1.4 Stopcock Grease. Acetone-insoluble, heat-stable silicone grease may be used, if necessary.

2.1.5 Temperature Gauge. Dial thermometer, or equivalent, to measure temperature of gas leaving impinger train to within 1°C (2°F).

2.1.6 Drying Tube. Tube packed with 6- to 16-mesh indicating-type silica gel, or equivalent, to dry the

FIG. 6-1 SO_2 SAMPLING TRAIN

gas sample and to protect the meter and pump. If the silica gel has been used previously, dry at 175°C (350°F) for 2 hours. New silica gel may be used as received. Alternatively, other types of desiccants (equivalent or better) may be used, subject to approval of the Administrator.

2.1.7 Valve. Needle valve, to regulate sample gas flow rate.

2.1.8 Pump. Leak-free diaphragm pump, or equivalent, to pull gas through the train. Install a small surge tank between the pump and rate meter to eliminate the pulsation effect of the diaphragm pump on the rotameter.

2.1.9 Rate Meter. Rotameter, or equivalent, capable of measuring flow rate to within 2 percent of the selected flow rate of about 1000 cc/min.

2.1.10 Volume Meter. Dry gas meter, sufficiently accurate to measure the sample volume within 2 percent, calibrated at the selected flow rate and conditions actually encountered during sampling, and equipped with a temperature gauge (dial thermometer, or equivalent) capable of measuring temperature to within 3°C (5.4°F).

2.1.11 Barometer. Mercury, aneroid, or other barometer capable of measuring atmospheric pressure to within 2.5 mm Hg (0.1 in. Hg). In many cases, the barometric reading may be obtained from a nearby national weather service station, in which case the station value (which is the absolute barometric pressure) shall be requested and an adjustment for elevation differences between the weather station and sampling point shall be applied at a rate of minus 2.5 mm Hg (0.1 in. Hg) per 30 m (100 ft) elevation increase or vice versa for elevation decrease.

2.1.12 Vacuum Gauge and Rotameter. At least 760 mm Hg (30 in. Hg) gauge and 0.40 cc/min rotameter, to be used for leak check of the sampling train.

2.2 Sample Recovery

2.2.1 Wash Bottles. Polyethylene or glass, 500 ml, two.

2.2.2 Storage Bottles. Polyethylene, 100 ml, to store impinger samples (one per sample).

2.3 Analysis

2.3.1 Pipettes. Volumetric type, 5-ml, 20-ml (one per sample), and 25-ml sizes.

2.3.2 Volumetric Flasks. 100-ml size (one per sample) and 1000-ml size.

2.3.3 Burettes. 5- and 50-ml sizes.

2.3.4 Erlenmeyer Flasks. 250 ml-size (one for each sample, blank, and standard).

2.3.5 Dropping Bottle. 125-ml size, to add indicator.

2.3.6 Graduated Cylinder. 100-ml size.

2.3.7 Spectrophotometer. To measure absorbance at 352 nanometers.

3. Reagents

Unless otherwise indicated, all reagents must conform to the specifications established by the Committee on Analytical Reagents of the American Chemical Society. Where such specifications are not available, use the best available grade.

3.1 Sampling.

3.1.1 Water. Deionized, distilled to conform to ASTM Specification D 1193-74, Type 3. At the option of the analyst, the KMnO_4 test for oxidizable organic matter may be omitted when high concentrations of organic matter are not expected to be present.

3.1.2 Isopropanol, 80 Percent. Mix 80 ml of isopropanol with 20 ml of deionized, distilled water. Check each lot of isopropanol for peroxide impurities as follows: shake 10 ml of isopropanol with 10 ml of freshly prepared 10 percent potassium iodide solution. Prepare a blank by similarly treating 10 ml of distilled water. After 1 minute, read the absorbance at 352 nanometers on a spectrophotometer. (Note: Use a 1-cm path length.) If absorbance exceeds 0.1, reject alcohol for use.

Peroxides may be removed from isopropanol by redistilling or by passage through a column of activated alumina; however, reagent grade isopropanol with suitably low peroxide levels may be obtained from commercial sources. Rejection of contaminated lots may, therefore, be a more efficient procedure.

3.1.3 Hydrogen Peroxide, 3 Percent. Dilute 30 percent hydrogen peroxide 1:9 (v/v) with deionized, distilled water (30 ml is needed per sample). Prepare fresh daily.

3.1.4 Potassium Iodide Solution, 10 Percent. Dissolve 10.0 grams KI in deionized, distilled water and dilute to 100 ml. Prepare when needed.

3.2 Sample Recovery

3.2.1 Water. Deionized, distilled, as in 3.1.1.

3.2.2 Isopropanol, 80 Percent. Mix 80 ml of isopropanol with 20 ml of deionized, distilled water.

3.3 Analysis

3.3.1 Water. Deionized, distilled, as in 3.1.1.

3.3.2 Isopropanol, 100 percent.

3.3.3 Thorin Indicator. 1-(o-arsonophenylazo)-2-naphthol-3, 6-disulfonic acid, disodium salt, or equivalent. Dissolve 0.20 g in 100 ml of deionized, distilled water.

3.3.4 Barium Perchlorate Solution, 0.0100 N. Dissolve 1.95 g of barium perchlorate trihydrate $[\text{Ba}(\text{ClO}_4)_2 \cdot 3\text{H}_2\text{O}]$ in 200 ml distilled water and dilute to 1 liter with isopropanol. Alternatively, 1.22 g of $[\text{BaCl}_2 \cdot 2\text{H}_2\text{O}]$ may be used instead of the perchlorate. Standardize as in Section 5.5.

3.3.5 Sulfuric Acid Standard, 0.0100 N. Purchase or standardize to ± 0.0002 N against 0.0100 N NaOH

which has previously been standardized against potassium acid phthalate (primary standard grade).

4. Procedure

4.1 Sampling

4.1.1 Preparation of Collection Train. Measure 15 ml of 80 percent isopropanol into the midget bubbler and 15 ml of 3 percent hydrogen peroxide into each of the first two midget impingers. Leave the final midget impinger dry. Assemble the train as shown in Fig. 6-1. Adjust probe heater to a temperature sufficient to prevent water condensation. Place crushed ice and water around the impingers.

4.1.2 Leak-Check Procedure. A leak check prior to the sampling run is optional; however, a leak check after the sampling run is mandatory. The leak-check procedure is as follows:

Temporarily attach a suitable (e.g., 0-40 cc/min) rotameter to the outlet of the dry gas meter and place a vacuum gauge at or near the probe inlet. Plug the probe inlet, pull a vacuum of at least 25 mm Hg (10 in. Hg), and note the flow rate as indicated by the rotameter. A leakage rate not in excess of 2 percent of the average sampling rate is acceptable. Note: Carefully release the probe inlet plug before turning off the pump.

It is suggested (not mandatory) that the pump be leak-checked separately, either prior to or after the sampling run. If done prior to the sampling run, the pump leak-check shall precede the leak check of the sampling train described immediately above; if done after the sampling run, the pump leak-check shall follow the train leak-check. To leak check the pump, proceed as follows: Disconnect the dry tube from the probe-impinger assembly. Place a vacuum gauge at the inlet to either the drying tube or the pump, pull a vacuum of 250 mm (10 in.) Hg, plug or pinch off the outlet of the flowmeter and then turn off the pump. The vacuum should remain stable for at least 30 seconds.

Other leak-check procedures may be used, subject to the approval of the Administrator, U.S. Environmental Protection Agency.

4.1.3 Sample Collection. Record the initial dry gas meter reading and barometric pressure. To begin sampling, position the tip of the probe at the sampling point, connect the probe to the bubbler, and start the pump. Adjust the sample flow to a constant rate of approximately 1.0 liter/min as indicated by the rotameter. Maintain this constant rate (± 10 percent) during the entire sampling run. Take readings (dry gas meter, temperatures at dry gas meter and at impinger outlet and rate meter) at least every 5 minutes. Add more ice during the run to keep the temperature of the gases leaving the last impinger at 20°C

(68°F) or less. At the conclusion of each run, turn off the pump, remove probe from the stack, and record the final readings. Conduct a leak check as in Section 4.1.2. (This leak check is mandatory.) If a leak is found, void the test run or use procedures acceptable to the Administrator to adjust the sample volume for leakage. Drain the ice bath, and purge the remaining part of the train by drawing clean ambient air through the system for 15 minutes at the sampling rate.

Clean ambient air can be provided by passing air through a charcoal filter or through an extra midget impinger with 15 ml of 3 percent H_2O_2 . The tester may opt to simply use ambient air, without purification.

4.2 Sample Recovery. Disconnect the impingers after purging. Discard the content of the midget bubbler. Pour the contents of the midget impingers into a leak-free polyethylene bottle for shipment. Rinse the three midget impingers and the connecting tubes with deionized, distilled water, and add the washings to the same storage container. Mark the fluid level. Seal and identify the sample container.

4.3 Sample Analysis. Note level of liquid in container, and confirm whether any sample was lost during shipment; note this on analytical data sheet. If a noticeable amount of leakage has occurred, either void the sample or use methods, subject to the approval of the Administrator, to correct the final results.

Transfer the contents of the storage container to a 100-ml volumetric flask and dilute to exactly 100 ml with deionized, distilled water. Pipette a 20-ml aliquot of this solution into a 250-ml Erlenmeyer flask, add 80 ml of 100 percent isopropanol and two to four drops of thiorin indicator, and titrate to a pink endpoint using 0.0100 N barium perchlorate. Repeat and average the titration volumes. Run a blank with each series of samples. Replicate titrations must agree within 1 percent or 0.2 ml, whichever is larger.

(Note: Protect the 0.0100 N barium perchlorate solution from evaporation at all times.)

5. Calibration

5.1 Metering System

5.1.1 Initial Calibration. Before its initial use in the field, first leak check the metering system (drying tube, needle valve, pump, rotameter, and dry gas meter) as follows: place a vacuum gauge at the inlet to the drying tube and pull a vacuum of 250 mm (10 in.) Hg; plug or pinch off the outlet of the flowmeter, and then turn off the pump. The vacuum shall remain stable for at least 30 seconds. Carefully release the vacuum gauge before releasing the flowmeter end.

ASME PERFORMANCE TEST CODES

Next, calibrate the metering system (at the sampling flow rate specified by the method) as follows: connect an appropriately sized wet test meter (e.g., 1 liter per revolution) to the inlet of the drying tube. Make three independent calibration runs, using at least five revolutions of the dry gas meter per run. Calculate the calibration factor, Y (wet test meter calibration volume divided by the dry gas meter volume, both volumes adjusted to the same reference temperature and pressure), for each run, and average the results. If any Y value deviates by more than 2 percent from the average, the metering system is unacceptable for use. Otherwise, use the average as the calibration factor for subsequent test runs.

5.1.2 Post-test Calibration Check. After each field test series, conduct a calibration check as in Section 5.1.1 above, except for the following variations: (a) the leak check is not to be conducted, (b) three or more revolutions of the dry gas meter may be used, and (c) only two independent runs need be made. If the calibration factor does not deviate by more than 5 percent from the initial calibration factor (determined in Section 5.1.1), then the dry gas meter volumes obtained during the test series are acceptable. If the calibration factor deviates by more than 5 percent, recalibrate the metering system as in Section 5.1.1, and for the calculations, use the calibration factor (initial or recalibration) that yields the lower gas volume for each test run.

5.2 Thermometers. Calibrate against mercury-in-glass thermometers.

5.3 Rotameter. The rotameter need not be calibrated, but should be cleaned and maintained according to the manufacturer's instruction.

5.4 Barometer. Calibrate against a mercury barometer.

5.5 Barium Perchlorate Solution. Standardize the barium perchlorate solution against 25 ml of standard sulfuric acid to which 100 ml of 100 percent isopropanol has been added.

6. Calculations

Carry out calculations, retaining at least one extra decimal figure beyond that of the acquired data. Round off figures after final calculation.

6.1 Nomenclature

C_{SO_2}	= concentration of sulfur dioxide, dry basis corrected to standard conditions, mg/dscm (lb/dscf)
N	= normality of barium perchlorate titrant, milliequivalents/ml
P_{bar}	= barometric pressure at the exit orifice of the dry gas meter, mm Hg (in. Hg)
P_{std}	= standard absolute pressure, 760 mm Hg (29.92 in. Hg)

T_m	= average dry gas meter absolute temperature, °K (°R)
T_{std}	= standard absolute temperature, 293°K (528°R)
V_a	= volume of sample aliquot titrated, ml
V_m	= dry gas volume as measured by the dry gas meter, dcm (dcf)
$V_{m(std)}$	= dry gas volume measured by the dry gas meter, corrected to standard conditions, dscm (dscf)
V_{soln}	= total volume of solution in which the sulfur dioxide sample is contained, 100 ml
V_t	= volume of barium perchlorate titrant used for the sample, ml (average of replicate rotations)
V_{tb}	= volume of barium perchlorate titrant used for the blank, ml
Y	= dry gas meter calibration factor
32.03	= equivalent weight of sulfur dioxide

6.2 Dry Sample Gas Volume, Corrected to Standard Conditions

$$V_{m(std)} = V_m Y \left(\frac{T_{std}}{T_m} \right) \left(\frac{P_{bar}}{P_{std}} \right) = K_1 Y \frac{V_m P_{bar}}{T_m} \quad (6-1)$$

where:

$$\begin{aligned} K_1 &= 0.3858 \text{ } ^\circ\text{K/mm Hg for metric units.} \\ &= 17.64 \text{ } ^\circ\text{R/in. Hg for English units.} \end{aligned}$$

6.3 Sulfur Dioxide Concentration

$$C_{SO_2} = K_2 \frac{(V_t - V_{tb}) N \left(\frac{V_{soln}}{V_a} \right)}{V_{m(std)}} \quad (6-2)$$

where:

$$\begin{aligned} K_2 &= 32.03 \text{ mg/meq for metric units.} \\ &= 7.061 \times 10^{-5} \text{ lb/meq for English units.} \end{aligned}$$

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6.01.5

**Brookhaven Controlled Condensation System
for the Measurement of Combustion Flue Gas Pollutants**

DETAILS OF THE METHOD**1. Scope**

1.1 Design. This method is primarily aimed at measuring the vapor phase flue gas content of sulfur dioxide (SO_2) and sulfuric acid (H_2SO_4) plus the metal sulfate (MSO_4) particulate suspended in the flue gas of combustion sources. The method also yields other gas phase constituents including water vapor (H_2O), oxygen (O_2), carbon monoxide (CO), carbon dioxide (CO_2), and nitrogen oxides (NO_x) as well as total suspended particulate (TSP) and the principal elemental composition of both the water soluble and insoluble portions of the particulate.

1.2 Application. Range of applicability of this method for SO_2 is 0.05 to 5000 ppm and for H_2SO_4 , 0.02 to 100 ppm (where ppm is parts-per-million on a molar or volume basis). The concentration of MSO_4 can be determined over the range 0.1 to 200 mg/m^3 as sulfate and TSP from 5 to 25,000 mg/m^3 . Water vapor is measured in the range from 5 to 15 percent, NO_x from 10 to 3000 ppm, and the other fixed gases from 0.005 to 15 percent. Principal elements are determined in both the insoluble and soluble portion of the TSP with a limit of detection of about 0.1 to 0.5 mg/m^3 .

1.3 Definitions

1.3.1 Total suspended particulate (TSP) is the weight of the total matter retained by the in-stack nozzle/filter assembly relative to the volume of flue gas sampled.

1.3.2 Sulfuric acid (H_2SO_4), which is generally a vapor at the usual temperature (300°F) of the flue gas, i.e., 160°C (320°F), but which can partially be present as an aerosol and partially as adsorbed on the particulate, is that amount of acid material found in the 100 percent isopropyl alcohol extract of the filter and the washings of the probe, controlled condensation coil, and back-up filter.

At flue gas temperatures exceeding 250°C , for example at an air heater inlet, the equilibrium between H_2SO_4 and SO_3 in the presence of H_2O can be shifted partially or entirely towards SO_3 . In this case temperature must be reduced to convert the SO_3 to H_2SO_4 in order to achieve efficient collection.

1.4 Significance. This method can be used for testing compliance with regulations governing the emissions of total suspended particulate (TSP) and oxides of nitrogen as well as plume visibility, which is dependent on TSP and sulfuric acid emission rate.

2. Summary of Method

2.1 Field Procedures. Flue gas is sampled isokinetically at nominally 10 L/min through an in-stack nozzle/filter assembly (NFA) which removes all suspended particulate but passes the H_2SO_4 vapor. Subsequently the H_2SO_4 is condensed to an aerosol and removed by impaction in the controlled condensation coil (CCC) and a back-up filter (BUF) of quartz fiber, both of which are maintained above the water dew point. A major portion of the flue gas water vapor is removed in an ice water jacketed condensation coil and receiver vessel, after which SO_2 is removed by oxidation to sulfate in dual impingers containing 3 percent H_2O_2 solution. The dried gas, a portion (0.5 L) of which is collected in a pre-evacuated bottle and a portion (0.5 L) in a pre-evacuated flask, is metered with a dry test meter at the exit of the pump. At the conclusion of the run, the NFA and BUF are placed in separate plastic bags and the probe, CCC, and two impingers are separately washed and the solutions retained for analysis. The volume of water collected is measured and then discarded, as it contains no significant amounts of SO_2 or NO_x .

2.2 Laboratory Procedures. The NFA is conditioned and weighed for TSP, washed with 100 percent isopropyl alcohol (IPA) to recover any adsorbed H_2SO_4 and with washing solution to recover MSO_4 , and conditioned and weighed again to obtain total insoluble suspended particulate. Any H_2SO_4 recovered from the NFA IPA wash is included with the total acid. The sulfate content of the water wash of the NFA is determined and principal elements are separately determined from the water wash and the insoluble particulate. Washings from the probe, CCC, and BUF together with the IPA wash of the NFA are used to determine total sulfuric acid. Sulfur dioxide is determined from the impingers, NO_x from the flask, and O_2 , CO , and CO_2 from the bottle.

3. Apparatus

3.1 Field Sampling Equipment. The equipment described in this section is either readily commercially available or easily fabricated. An overall layout of the equipment (cf., Fig. 1) shows the sampling arrangement of the NFA, probe and H_2SO_4 controlled condenser coil (CCC) with aluminum probe assembly support, BUF, water condenser and receiver, SO_2 impingers, dryer, pump, evacuated bottle, NO_x flask, and dry test meter.

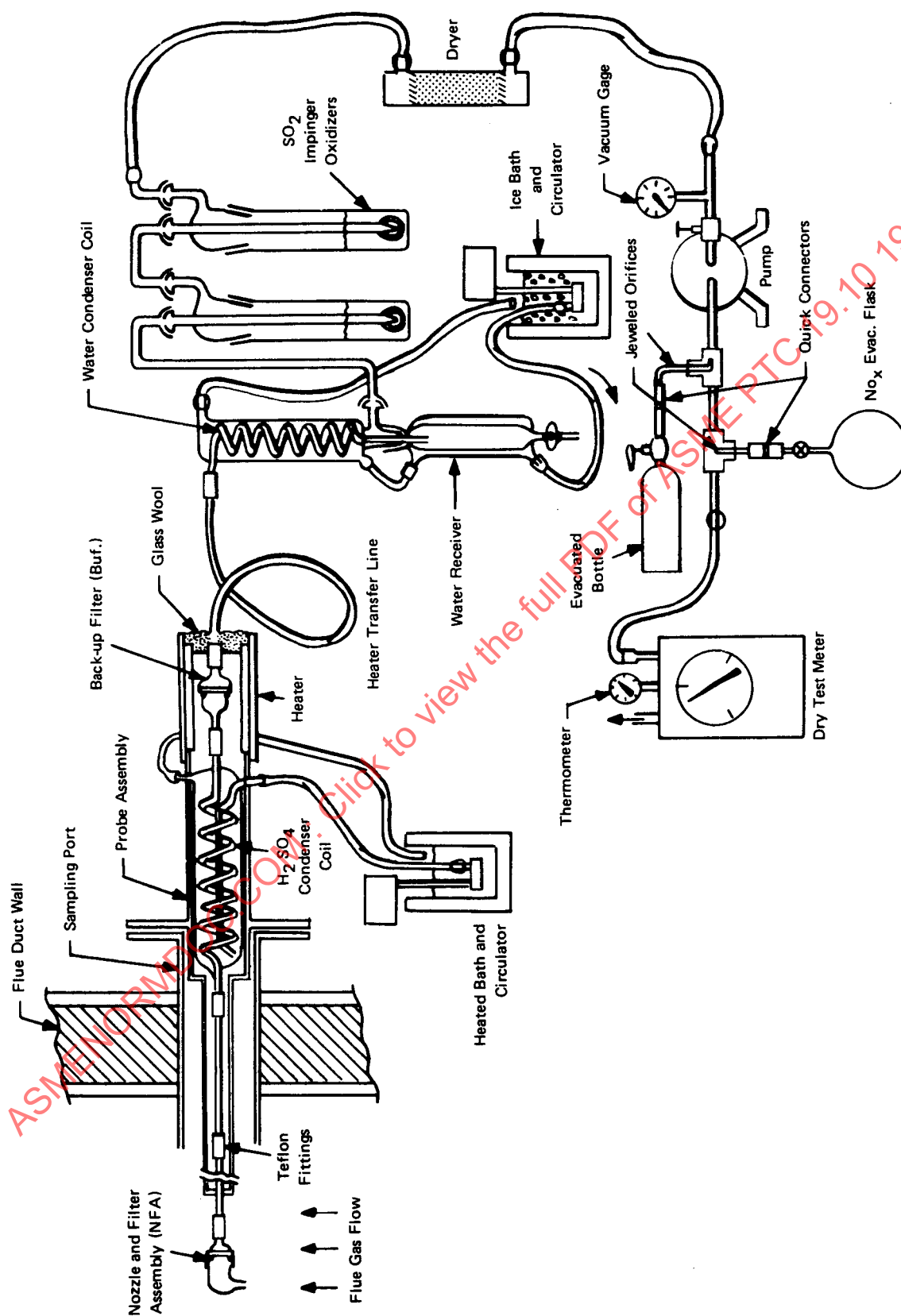


FIG. 1 SCHEMATIC DIAGRAM OF BNL CONTROLLED CONDENSATION SCHEME

3.1.1 Nozzle/Filter Assembly (NFA)

One end of the assembly is fabricated from a 60 mL coarse fritted Pyrex Buchner funnel as shown in Fig. 2. On the face of the frit, a 40 mm diameter Pallflex tissue quartz fiber filter (QAO 2500) that has been phosphoric acid washed and heat treated to remove organic binder (Reference 1) is held in place with a Vitron O-ring (satisfactory up to 250° to 300°C). The nozzle half of the assembly makes a leak tight seal against the O-ring when held with the four stainless steel spring retainers. The assembled nozzles are shown in Fig. 3. The smaller nozzle (0.37 cm diameter) is used for sampling where flue gas velocities are 100 ft/sec and the larger (0.79 cm diameter) for velocities near 20 ft/sec. A 3/8-in. Teflon compression fitting secures the NFA to the end of the glass probe and the entire NFA is wired to the aluminum probe sheath to prevent loss during removal from the sampling port.

3.1.2 Probe Assembly

Borosilicate glass tubes, 3/8-in. in outside diameter and in 3-ft long sections, are interconnected with Teflon compression fittings and slide into a nominal 1-in. aluminum pipe sheathing which is fashioned to the appropriate length from 1-, 2-, and 4-ft sections. Only the 3-ft glass section nearest the H₂SO₄ condenser coil (cf., Fig. 1) is wrapped with a self-sticking heater tape to maintain the probe temperature at about 160° to 170°C (320° to 340°F) in the region passing through the flue duct wall; the balance of the probe as well as the NFA, is at flue gas temperature. The end of the aluminum probe in the flue gas has a pipe cap with a 3/8-in. hole to allow the passage of the glass probe for connection to the NFA. The other end is connected directly to the H₂SO₄ condenser coil aluminum housing which is a 12-in. length of nominal 2½-in. pipe with a 6½-in. flange.

3.1.3 H₂SO₄ Controlled Condenser Coil (CCC)

The coil consists of an outer Pyrex jacket about 2 in. in diameter around a 17-turn coil of 6 mm I.D. Pyrex tubing. Provision is made to control the temperature of the jacket at 60°C (140°F) with water from a heated bath and circulator.

3.1.4 Back-up Filter (BUF)

Although most of the H₂SO₄ is removed in the CCC, some of the aerosol would escape collection if not retained by the heated BUF, located immediately after the CCC as shown in Fig. 1. The BUF, which is essentially identical to the NFA except for straight 3/8-in. O.D. tubulations on either end, is also maintained at 60°C (140°F) by an electrically controlled clam-shell heater. Tests have demonstrated better than 98 percent efficiency for the collection of H₂SO₄ aerosol. (Reference 2.)

3.1.5 Water Condenser and Receiver

A 13-turn jacketed condenser, similar in dimensions to the CCC, and a 150-mL jacketed cylinder, each cooled to

about 0.5°C with circulating ice water, are connected to the BUF through an electrically heated Teflon transfer line. Polyvinyl chloride (PVC) tubing is used to connect the circulators to the jacketed devices and also for a number of subsequent sampling train interconnections as indicated in Fig. 1.

3.1.6 SO₂ Impingers

Two Smith-Greenburg 500 mL impingers containing H₂O₂ solutions are used to oxidize and remove SO₂.

3.1.7 Dryer

A total of about 400 mL of Drierite is used to remove remaining moisture to prevent corrosion of the pump and dry test meter. The drying agent is divided equally between two tubes, with one being replaced as soon as the second shows signs of collecting water (e.g., on color change).

3.1.8 Pump

The rotary vane compressor is capable of delivering 10 L/min while pulling on a vacuum of up to 20 in. Hg. Flow rate during sampling is adjusted by the valve on the leak-tight pump.

3.1.9 Evacuated Sample Bottle

A 1-L steel sampling bottle equipped with a toggle valve, quick connect, and 20 mL/min orifice with filter is teed into the exhaust of the pump.

3.1.10 NO_x Flask

A pre-evacuated 1000 mL glass flask containing acidified peroxide solution and equipped with a glass valve and 20 mL/min critical orifice is teed into the exhaust line after the evacuated bottle.

3.1.11 Dry Test Meter

With a resolution of 1 L per revolution, total volume of sample collected is recorded by the dry test meter. Using a stop watch, sampling rates are determined.

3.1.12 Plastic Bottles and Bags

Polyethylene bottles are used to retain the SO₂ impinger solutions and the wash solutions from the probe and the CCC. Polyethylene zip-lock type bags are used to package the NFA and BUF for subsequent analytical work up.

3.1.13 Thermometer

Dial thermometers are used to measure the temperature of the heated bath, ice bath, and dry test meter. A multiprobe thermocouple meter is used to record temperatures of the flue gas and the heaters for the probe, BUF, and Teflon transfer line.

3.1.14 Ambient Thermometer/Humidity

A recording thermometer and relative humidity indicator is necessary for accurate moisture material balances.

3.1.15 Pitot Tube and Manometer

Either an inclined manometer or pressure gage with 0.01-in. H₂O sensitivity is needed to measure flue gas velocity as detected by a Pitot tube.

INSTRUMENTS AND APPARATUS



FIG. 2

3.1.16 Oxygen Analyzer

Furnace oxygen content is obtained by sampling and recording with a continuous electrochemical analyzer, drawing a sample through a water condenser and dryer to prevent the formation of liquid water. The measurement is generally performed at the air heater inlet since air leakage into the ducting is usually minimal up to that point. Oxygen determinations at locations further downstream, including the sulfur oxides sampling locations, are always contaminated by air leakage and are not a reliable measure of the furnace oxygen or excess combustion air. A controller sequentially provides for sampling, zeroing, and spanning (3 percent by volume of O_2) the analyzer over a 20-min cycle time.

3.2 Laboratory. The equipment described in this section is standard laboratory apparatus and instrumentation.

3.2.1 Analytical Balance

An automatic balance with 0.1 mg resolution and a capacity of about 100 g is required.

3.2.2 Volumetric Glassware

Pipettes, burettes, cylinders, flasks, beakers, and funnels. Titrations are performed with a 5 mL burette having 0.01 mL divisions.

3.2.3 Vacuum Oven

Having temperature control up to 100°C under vacuum is necessary for drying certain samples.

3.2.4 Mortar with Pestle

To obtain a homogeneous sample of the particulate from the filter, the entire filter is ground in a mortar.

3.2.5 pH Meter

Titration of H_2SO_4 washed from various parts of the sampling apparatus requires a pH meter with a resolution of 0.05 pH units.

3.2.6 Ion Chromatograph

Determination of total water soluble metal sulfates and the NO_x concentration requires the ion chromatograph as the best choice for determination of sulfate and nitrate ions.

3.2.7 Atomic Absorption Spectrometer

Principal elements in the flue gas particulate are determined by atomic absorption spectroscopy.

3.2.8 Total Carbon Analyzer

Soot or carbon level in the fly ash is determined from the carbon on the in-stack filter by conversion to CO_2 for detection by thermal conductivity in a CHN elemental analyzer.

3.2.9 Shaker

Proper reaction of the NO_x flue gas sample with the oxidizing solution requires an agitator or shaker.

3.2.10 Thermal Conductivity Gas Chromatograph

The gas chromatograph with a vacuum pump for sample transfer from the partially evacuated steel bottles is needed to determine sampling location O_2 , CO , and

CO_2 . A vacuum gage (0-800 mm Hg) is needed for measuring sample volume.

4. Reagents

Standard reagent grade chemicals are used in all phases of sampling and analysis.

4.1 Water. Deionized water with a sulfate content of less than 0.05 g/mL is used.

4.2 Isopropyl Alcohol. Solutions of 100 percent IPA and a solution of 5 percent IPA in water (washing solution) are used.

4.3 Sulfuric Acid (0.05M). Volumetric prepared standards are used.

4.4 Sodium Hydroxide (0.1 and 0.02 M). Volumetric prepared standards are used.

4.5 Buffer Solutions (pH 4 and 7). Prepared buffer solutions are used to span the pH meter.

4.6 Hydrogen Peroxide (3 Percent). Fresh solutions are prepared from 30 percent H_2O_2 reagent as needed for the SO_2 impingers and the NO_x flask.

4.7 Oxygen Gas Standard (3 Percent by Volume). A prepared O_2 standard is required to certify the span of the O_2 field analyzer.

4.8 Sulfate and Nitrate Solutions. The ion chromatograph is calibrated for sulfate from 0.5 to 50 $\mu\text{g/mL}$ and for nitrate from 0.1 to 50 $\mu\text{g/mL}$.

4.9 Cation Standard Solutions. The atomic absorption spectrometer is calibrated for principal metals using standard cation solutions.

5. Sampling Preparation

Before field sampling can commence, certain preparations are necessary in the laboratory.

5.1 Nozzle and Filter Assembly (NFA). Each field sample is performed with a separate NFA that has been assembled, conditioned by drawing 15 to 20 L of room air through each, and weighed. Record this tare weight to within ± 0.1 mg. Each NFA has a numbered and lettered code for identification and is packed in a separate polyethylene zip-lock bag.

5.2 Glassware. All glassware including probes, controlled condensation coil, back-up filters, and SO_2 impingers are clean and free of any trace acid.

5.3 Back-up Filter (BUF). A separately labeled and assembled BUF is placed in a separate polyethylene zip-lock bag, one for each sampling run to be performed.

5.4 SO_2 Impinger Solution. For each run, two separate 250-mL polyethylene bottles containing 150 mL each of the 3 percent H_2O_2 solution are needed for the SO_2 impingers.

5.5 Evacuated Bottle. Each steel bottle is evacuated and closed by the toggle valve prior to deployment.

5.6 NO_x Evacuated Flask. Place 10 mL of 0.05 M H₂SO₄ and 10 drops of 3 percent H₂O₂ into each flask, and quickly (30s) evacuate to a residual pressure of about 10 mm Hg. Close the vacuum source, allow 1 to 2 min for the pressure in the flask to equilibrate, record the temperature and the pressure and then close the valve on the flask. (Note: Testing is being performed to determine if 0.02 M H₂SO₄ can be used to minimize the slight interference on the ion chromatograph, since the generally small nitrate peak, about 2 to 10 µg/mL, just precedes the larger sulfate peak, about 480 µg/mL; the volume of the NO_x flask should be precisely known to within ±1 mL.)

6. Field Sampling

6.1 Preliminary. Crushed ice is required for the water condenser circulator bath. A notebook is required to record pertinent data, a stopwatch for determining flow rates, and several extension cords for power. Set up the portable recording temperature and humidity device in a shady location.

6.2 Oxygen Analyzer. Install the portable oxygen analyzer and sequential controller at a sampling port near the inlet of the air heater. Traverse the depth of the duct at the port located near the center of the face of the duct to establish a location that provides a representative sample. Be especially careful to exclude leaks where the probe attaches to the port. Attach the oxygen analyzer to a recorder.

6.3 Nozzle Size Selection. Measure the velocity in the duct to be sampled using the standard Pitot tube and manometer. Velocity is calculated from

$$V, \text{ ft/sec} = 3.90 \sqrt{\Delta P (t + 273.2)} \quad (1)$$

where ΔP is the manometer reading in inches of water and t is the temperature of the flue gas (°C). Based on the velocity, a nozzle size is selected, such that isokinetic sampling can be performed with a flow rate in a range of from 4 to 6 L/min if total particulate is in the range of 0.5 to 2.5 g/m³ (coal-fired ESP inlet), 8 to 10 L/min if total particulate is from 200 to 1000 mg/m³ (coal-fired ESP outlet), and 10 to 14 L/min if total particulate is less than 200 mg/m³ (oil-fired unit). Table 1 shows that three nozzle sizes from 3.5 mm to 8 mm would provide the necessary range to cover velocities expected at conventional combustion sources. (Note: The efficiency of the back-up filter (BUF) for collection of H₂SO₄ aerosol breaking through the controlled condensation coil (CCC) has been shown to be greater than 98 percent removal at flow rates from 0.5 to 15 L/min² (Reference 2). The efficiency of the SO₂ impinger was found to be 95 to 98 percent for one impinger and greater than 99 percent for two in series at flow rates up to 15 L/min³ (Reference 3)).

TABLE 1

Nozzle Diameter and Sampling Flow Rate Selection
Flue Gas Temperature of 300°F (149°C)

ΔP , in. of water	Flue Gas Velocity, ft/sec	Flow Rate ^a , L/min at 16°C (60°F)		
		3.5 mm ^b	5.5 mm ^b	8 mm ^b
0.02	11	...	...	6.4
0.05	18	...	...	10.2
0.10	25	...	6.8	14.4
0.20	36	...	9.6	...
0.4	51	5.5	13.6	...
1.0	80	8.7	...	...
1.6	101	11.0	...	...
2.4	124	13.5	...	...

^a Flow rate = $4.40 A \sqrt{\Delta P \times (t + 273.2)}$; cf., Section 6.3 for definition of terms.

^b Nozzle diameter.

6.4 System Assembly. Assemble appropriate lengths of glass probes with Teflon fittings and connect to the H₂SO₄ condenser coil with a Teflon fitting. After assembling the appropriate length of aluminum probe shield, slide the glass probe and condenser into the probe assembly, taking care that the connectors for the heated bath tubing slide into the appropriate slots. Use a weather stripping dough to make a leak-tight seal between the probe assembly and the end of the H₂SO₄ condenser, otherwise air leakage could occur between the condenser and the probe assembly when sampling at subatmospheric ducts, resulting in undesirable cooling of the probe.

Connect the BUF and its supported heating mantle to the back end of the H₂SO₄ condenser coil. Next, connect the appropriately sized NFA at the end of the probe using a Teflon connector and secure with wiring to prevent loss, especially when removing the assembly from the sampling port. Finally, attach the thermocouple lead near the filter in order to record flue gas temperature.

On a ring stand or small portable rack, appropriately interconnect the water condenser and receiver vessel with the two SO₂ impingers. Fill each impinger with 150 mL of the 3 percent H₂O₂ solution. Appropriately interconnect the impingers with the dryer and pump as shown in Fig. 1, using PVC tubing. Finally, connect the evacuated bottle, the NO_x flask, and the dry test meter. Record in the notebook, the NFA, BUF, NO_x flask, and evacuated bottle identifying numbers.

Before installing the probe assembly into the sampling port, a vacuum leak check is essential. Temporarily connect the heated transfer line between the BUF and the water condenser, plug the end of the NFA (e.g., someone can cover the inlet with his finger), and turn the pump on.

The flow of air, as indicated by the dry test meter, should decrease to essentially zero. *Caution:* remove plug or finger from the NFA before turning the pump off, otherwise part of the solution in the impingers would blow backwards out the impinger inlet. Similarly, the filters could be dislodged from their holders by the reverse flow of air.

When the system has been determined to be leak-tight, disconnect the heated transfer line from the BUF and install the entire probe assembly into the sampling port, making certain that the nozzle is facing *downstream*, that is, in the direction of flow. Using C-clamps, secure the probe assembly flange to the sampling port flange. Reconnect the heated transfer line and start the flow of water from the heated bath and ice bath. The heated bath should be adjusted to maintain the condenser water exit temperature at 58° to 62°C (136° to 144°F). At the same time adjust the temperature of the 3-ft heated probe section to about 160°C (320°F), and the BUF and heated transfer line to about 60°C (140°F). Set the dry test meter dials to zero.

6.5 Sampling. With all systems at the appropriate temperatures, rotate the probe assembly 180 deg so that the nozzle is facing upstream into the direction of flow, start the pump, and begin timing for a sample duration of 30 min. Adjust the valve on the pump to achieve the desired number of liters per minute as determined earlier (use the dry test meter and a watch). Record the pump vacuum and dry test meter temperature and, after 3 or 4 min, open the valves on the evacuated bottle and the NO_x flask.

Maintain a check on the temperature of the flue gas, probe, heated bath, BUF, and ice bath. Add ice to the bath as needed. Routinely check the sampling rate, as particulate building up on the filter in the NFA will increase the resistance to flow. To maintain the desired flow rate, the pump vacuum will have to be increased gradually during the 30 min sampling period by opening the pump valve. If pump vacuum must be increased to above 20 in. Hg vacuum in order to maintain flow (due to high particulate levels), terminate the run earlier than 30 min. Table 2 lists data to be recorded.

About 1 to 2 min before the end of the 30 min sampling period, close the valves on the evacuated bottle and the NO_x flask. Record the final pump vacuum.

At exactly 30 min, close the pump valve, turn off the pump, and rotate the probe assembly such that the nozzle faces downstream. It is important that the pump valve be closed first to prevent reverse flow of air which would blow out the impinger solution and also dislodge the filters. Record the elapsed sampling time and the volume indicated on the dry test meter. Disconnect the heated transfer line, turn off all heaters, remove the probe assembly, and allow to cool. Drain the water from the water receiver into

TABLE 2

Sampling Parameters Recorded

1. Run number	13. Pump vacuum: Start and finish
2. Location	14. Temperatures:
3. Date	Stack
4. Time	Probe
5. Probe length	Condenser
6. NFA number	BUF
7. BUF number	Transfer line
8. Total run time	Ice bath
9. Barometric pressure	15. Evacuated bottle number
10. Ambient temp. and humidity	16. NO _x flask number
11. Dry test meter volume	17. Pitot tube pressure reading
12. Dry test meter temperature	18. Water condenser volume

Unit Operating Conditions

1. Furnace O₂
2. Power level
3. Fuel composition
4. Electrostatic precipitator conditions
5. Furnace CO

a volumetric cylinder and record the volume of water.

Discard the water. (Note: The ice bath condenser water has been shown to contain artifact sulfuric acid, form from the oxidation of a slight amount of dissolved SO₂ from the flue gas (Reference 1). The acidity in this water must not be included in the H₂SO₄ measurement. The amount of SO₂ and NO_x retained in the ice water is a negligible fraction of that collected, respectively, in the SO₂ peroxide impingers and the NO_x flask can be disregarded.)

Pour the contents of each SO₂ impinger into separate 250-mL labeled polyethylene bottles, rinse the impingers, and include the washings in the respective bottles. Disconnect the evacuated bottle and the NO_x flask and store for return to the laboratory.

Remove the NFA and the BUF and store in their respectively labeled zip-lock polyethylene bags. Slide the probe and H₂SO₄ condenser coil out of the probe assembly. Disconnect the coil from the probe, rinse each with washing solution (5 percent isopropyl alcohol in water), and store the washings in separately labeled polyethylene bottles for subsequent analysis of the dissolved H₂SO₄. The system is now ready for reassembly for another sampling run.

7. Analysis of Field Samples

7.1 Nozzle/Filter Assembly (NFA). The NFA contains some adsorbed H₂SO₄ plus all the particulate matter from

the flue gas that was sampled. From the NFA, the total suspended particulate (TSP), the soluble sulfates, the principal elements in the soluble and insoluble fraction, and the carbon content of the particulate will be determined.

7.1.1 Cleaning and Weighing

Before the NFA can be weighed for gravimetric determination of TSP, any external material adhering to the outside of the assembly must be removed. Connect a rubber tube between the inlet and outlet of the NFA to prevent water from entering the filter during the washing and carefully flush the outside surfaces under running tap water to remove all particulate. Remove the rubber tube and place the NFA in a vacuum oven at 65°C for an overnight (16 hr) period. Maintain a beaker of water in the vacuum oven to limit the extent of drying of the filter, thereby minimizing the chances of loss of adsorbed H_2SO_4 or of converting soluble vanadyl sulfate pentahydrate into the insoluble form. However, drying does remove any residual adsorbed SO_2 if present. Following vacuum drying, condition the NFA by drawing 15 to 20 L of room air through each in a laminar flow hood. The weight of the NFAs can then be recorded to within ± 0.1 mg.

7.1.2 Recovery of Adsorbed H_2SO_4

Flush 20 to 25 mL of 100 percent isopropyl alcohol (IPA) through the NFA (squirt the IPA into the nozzle end) to remove H_2SO_4 that has adsorbed onto the glass surfaces of the NFA. The amount of acid found on the NFA is usually 5 percent or less of the total acid recovered. The 100 percent IPA does not dissolve any metal sulfates. Dilute to about 100 mL with H_2O and titrate with 0.02 M NaOH to the same pH of a similar IPA blank solution (generally a pH of 5.6). Record the milliequivalent of acid found.

7.1.3 Recovery of Soluble Particulate

Wash the NFA with three 10-mL aliquots of water through the filter into a vacuum Erlenmeyer flask, drawing the water through at a slow rate to avoid physical removal of insoluble material. Dilute to 50 mL with H_2O .

7.1.3a Determination of Soluble Sulfate. A portion of the above solution is injected into an ion chromatograph for determination of sulfate content. Record the μg of sulfate per mL of solution (i.e., ppm) found.

7.1.3b Determination of Principal Metals. The balance of the solution is used to determine the principal water soluble elements by using an atomic absorption spectrometer. At oil-fired units the elements are generally Mg, V, Na, Ca, Ni, and Fe. At coal-fired units the same metals can be found and, in addition, K is a principal soluble metal. All of the water soluble metals in solution have been shown to be present originally as sulfates. Record the μg of metal per mL of solution found.

7.1.4 Reweighing of NFA

Again condition the NFA by drying overnight in the 65°C vacuum oven containing a beaker of H_2O and then, by pulling several liters of room air through the assembly. Record the weight to within ± 0.1 mg. The difference in weight between that found with all the particulate and that found after washing, can be related to the total weight of water soluble material.

7.1.5 Recovery of Insoluble Particulate

Carefully disassemble the NFA, remove the filter, and grind to a powder in a mortar with pestle.

7.1.5a Determination of Principal Metals. Decompose a representative sample of the ground filter in a crucible by a low temperature ignition in a 450°C oven for 1 hr to remove any organic matter. Then dissolve in 10 percent HCl containing a few drops of concentrated HNO_3 and dilute to 25 mL. Determine the principal metals as in Section 7.1.3b. For oil-fired units, the principal metals are Mg, V, Na, Ni, Fe, and Ca. At coal-fired units, K and Al are also determined. The high silicon content present in coal ash can be determined only if the sample is dissolved in HF and if a suitable correction can be made for the filter, which is 100 percent silica. Record concentrations as mg/g of sample.

7.1.5b Determination of Carbon. A portion of the ground filter which, of course, has not been ignited, but has been dried further in a low temperature oven, is placed in the sample boat of the elemental analyzer for determination of the carbon content by conversion to CO_2 . Record the carbon content as mg C per g of sample.

7.2 Determination of H_2SO_4 . During field sampling, sulfuric acid will have been retained primarily by the H_2SO_4 condenser coil, but a significant amount is found in the probe and the BUF, with a trace in the NFA (cf., Section 7.1.2).

7.2.1 Probe and CCC H_2SO_4

Bring the separately collected washings of the probe and controlled condensation coil (CCC) each to a volume of about 50 mL and titrate with 0.02 M NaOH from the 5 mL burette, using the pH meter to determine the end point (generally a pH of 5.6). Multiply the volume of titrant (measured to within 0.005 mL) in each case by the normality of the NaOH (0.02 N), and record the milliequivalents of acid found in both the probe and CCC washings (record to the nearest 0.0001 meq).

7.2.2 BUF H_2SO_4

Flush three 15 mL aliquots of washing solution (5 percent IPA in water) through the inlet end of the back-up filter (BUF), making certain to contact all of the inside surfaces. Combine the washings, titrate with 0.02 M NaOH as in Section 7.2.1 above, and record the milliequivalents of H_2SO_4 .

7.2.3 Total H₂SO₄

The sum of the milliequivalents of H₂SO₄ found in the NFA (cf., Section 7.1.2), the probe and the CCC (Section 7.2.1), and the BUF (Section 7.2.2) represents the total milliequivalents of sulfuric acid to be recorded. (Note: In the absence of H₂SO₄ but with SO₂ present, the measured amount of acidity is always negligible, i.e., less than 0.0001 meq (Reference 2)).

7.3 Determination of SO₂. Bring the contents of the respective SO₂ impinger solutions each to a volume of 250 mL. Dilute a 2-mL aliquot of the first impinger solution and a 25-mL aliquot of the second, each of 50 mL, and titrate to the blank solution end point (pH ~ 5.6) using 0.1 M NaOH. Multiply the volume of titrant for the first impinger solution by 12.5 and add to it the volume of titrant from the second impinger titration. Record this number as the total milliequivalents of SO₂.

7.4 Analysis of Evacuated Bottle. Connect the bottle to the sample inlet port of the thermal conductivity gas chromatograph sample valve containing an 8-mL sample loop and evacuate up to the valve on the bottle. When a vacuum has been attained as indicated by a zero reading on a 0-800 mm Hg vacuum gage, close the vacuum valve, and open the sample bottle. Record the pressure and temperature and inject the sample into a molecular sieve column for separation of the oxygen, nitrogen, carbon monoxide, and CO₂ (Reference 4). Record the integrator areas for each constituent. Calibrate the chromatograph using room air at ambient conditions in a 0.5-mL sample loop, recording temperature and barometric pressure as well as peak areas.

7.5 Analysis of NO_x Flask. Shake the NO_x flask overnight on a laboratory agitator in order to ensure complex oxidation of the NO into nitrate. Then connect the flask to a vacuum gage (e.g., a Wallace-Tiernan), reduce the pressure in the line up to the valve on the flask to about 400 mm Hg, open the flask valve, and record the pressure and the temperature. Remove the flask, pour out the solution, and rinse several times. Dilute to 50-mL and determine nitrate by a calibrated ion chromatograph following standard procedures (Reference 5). Record concentration as μg NO₃⁻ per mL.

8. Calculation of Results

This section presents the basic equations necessary to perform the desired calculations.

8.1 Predicted SO₂ and H₂O. Based on the composition of the fuel and the amount of excess air, a stoichiometrically balanced equation can be derived for the combustion process. A typical case for a fuel oil is shown in Table 3. Equation (2) in the table defines the total number of moles of flue gas, F , from the combustion of

TABLE 3

Calculations for Predicted Flue Gas Concentrations

STOICHIOMETRIC EQUATION

Typical Fuel Oil Composition (2.3 percent S):

	Weight Fraction	Moles	Moles per Unit C	Moles O ₂ Required
C	0.8610	0.07168	1.000	1.000
H	0.1120	0.1111	1.550	0.3875
N	0.0030	0.00021	0.0030	0.0015
S	0.0230	0.00072	0.0100	0.0100
ash	0.0009			1.399 × f

where $f = 1 + \text{fraction excess air}$

Total Air Components (moles per mole of fuel):

$$1.399 f \text{ O}_2 + 5.215 f \text{ N}_2 + 0.0624 f \text{ A} + 0.0022 f \text{ CO}_2 = 6.679 f$$

H₂O from Air Humidity:

$$\frac{P_{\text{H}_2\text{O}}}{760 - P_{\text{H}_2\text{O}}} \times 6.679 f = W, \text{ moles H}_2\text{O/mole fuel} \quad (1)$$

$$\begin{aligned} & \text{CH}_{1.55}\text{N}_{0.003}\text{S}_{0.01} + 1.399 f \text{ O}_2 + 5.215 f \text{ N}_2 + 0.0022 f \text{ CO}_2 \\ & + W \text{ H}_2\text{O} + 0.0624 f \text{ A} \\ & \rightarrow 1.0024 \text{ CO}_2 + 1.399 (f-1) \text{ O}_2 + 5.215 f \text{ N}_2 + (0.775 + W) \text{ H}_2\text{O} \\ & + 0.010 \text{ SO}_2 + 0.0624 f \text{ A} \end{aligned}$$

TOTAL FLUE GAS MOLES

$$F = 0.3884 + 6.6764 f E W \quad (2)$$

EXCESS AIR

$$[\text{O}_2]_g = \frac{139.9 (f-1)}{F}$$

where:

$$[\text{O}_2]_g = \text{percent excess O}_2 \text{ in furnace}$$

$$f = 1 + \text{fraction excess air}$$

$$f = \frac{139.9 + (0.3884 + W) [\text{O}_2]_g}{139.9 - 6.6764 [\text{O}_2]_g} \quad (3)$$

SULFUR DIOXIDE

$$\text{SO}_2, \text{ ppm} = \frac{10^4 (\text{wt } \% \text{ S in fuel})}{2.3 F} \quad (4)$$

WATER VAPOR

$$\text{H}_2\text{O}, \% = \frac{(0.775 + W) 100}{F} \quad (5)$$

one mole of carbon in the fuel. The unknowns in equation (2) can be computed from equation (1) and (3). By substituting equation (1) for W in equation (3), f can be calculated explicitly in terms of the partial pressure of ambient humidity and furnace oxygen level, $[O_2]$, which was determined from the portable analyzer.

Once F has been calculated, the predicted levels of SO_2 and H_2O are calculated from equations (4) and (5), respectively. The benefit of calculating the expected levels of SO_2 and H_2O is that they can be compared to the corrected measured concentrations. If the agreement is good (i.e., within ± 5 percent), a certain degree of confidence can be placed in the representativeness of the sample.

8.2 Measured SO_2 , H_2O , Sulfate, and H_2SO_4 . First, the volume of flue gas sampled must be calculated. The volume of the dried flue gas, V_{FD} , as measured by the dry test meter is given by

$$V_{FD} = M.R. \times \frac{B.P.}{760} \times \frac{288.7}{(t + 273.2)} \quad (2)$$

where:

V_{FD} = dry flue gas volume, L (NTP),

$M.R.$ = dry test meter reading, L

$B.P.$ = barometric pressure, mm Hg, and

t = meter temperature, $^{\circ}C$.

(Note: NTP is $60^{\circ}F$ ($15.6^{\circ}C$) and 1 atm)

A portion of the original flue gas contained water vapor which was not seen by the dry test meter since most was condensed out and the balance was removed by the dryer. That portion which was removed by condensation is given by

$$V_{WC} = \frac{V_{WL}}{18} \times 23.7 \quad (3)$$

where:

V_{WC} = vapor volume of condensed H_2O , L (NTP), and

V_{WL} = volume of condensed H_2O , mL.

That portion of water which was removed by the dryer is given by

$$V_{WA} = \frac{P_o}{P_v - P_o} \times V_{FD} \quad (4)$$

where:

V_{WA} = vapor volume of adsorbed H_2O , L (NTP),

P_o = vapor pressure of H_2O at ice bath temp., mm Hg, and

P_v = vacuum pump absolute inlet pressure, mm Hg.

The sum of equations (2), (3), and (4) is the volume of flue gas containing the water vapor, that is,

$$V_{FW} = V_{FD} + V_{WC} + V_{WA} \quad (5)$$

From the volume of the wet flue gas, V_{FW} , and the milliequivalents of SO_2 and H_2SO_4 , the measured flue gas concentration of those constituents can be calculated:

$$C_m = 11845 \times \frac{MEQ}{V_{FW}} \quad (6)$$

where:

C_m = measured concentration of SO_2 or H_2SO_4 , volume ppm,

MEQ = milliequivalents of SO_2 or H_2SO_4 , and

V_{FW} = volume of wet flue gas, L (NTP).

The measured concentration of H_2O in the flue gas is simply given by

$$C_{H_2O} = \frac{(V_{WC} + V_{WA})}{V_{FW}} \times 100 \quad (7)$$

where:

C_{H_2O} = measured concentration of H_2O , volume percent.

From the concentration of sulfate found in the 50 mL of filter wash solution, the concentration in the flue gas is given by

$$C_{SO_4} = 12.34 \times \frac{SO_4}{V_{FW}} \quad (8)$$

where:

C_{SO_4} = measured concentration of flue gas sulfate, vol ppm, and

SO_4 = concentration of sulfate in 50 mL solution, wt ppm.

(Note: Multiplying C_{SO_4} by 4.055 gives sulfate as mg SO_4 / m^3 .)

8.3 Measured Oxygen at Sampling Location. By taking into account the higher O_2 content in the sampled flue gas compared to that present in the furnace region, correction can be made for dilution of the flue gas from air leaks in the ducting. The O_2 in the sampled flue gas is determined from the evacuated bottle sample as analyzed on the gas chromatograph.

The O_2 calibration factor of the chromatograph was calculated by

$$C_F = \frac{V_L}{A} \times \frac{P - P_{H_2O}}{760} \times \frac{273.2}{(t + 273.2)} \times 0.2188 \quad (9)$$

where:

C_F = calibration factor,

V_L = volume of sample loop, mL,

A = integrator area,

P = pressure of sample, mm Hg,
 P_{H_2O} = partial pressure of humidity, mm Hg, and
 t = temperature of sample loop, °C
 for samples of lab air used in the calibration oxygen.
 The concentration of oxygen actually found in the bottle is given by

$$C_{O_2} = \frac{100 \times A \times C_F}{V_L} \times \frac{760}{P} \times \frac{(t + 273.2)}{273.2} \times \frac{V_{FD}}{V_{FW}} - \frac{6.24f}{F} \quad (10)$$

where:

C_{O_2} = measured flue gas O_2 , vol percent.

The last term in equation (10) is a correction for the amount of argon in the flue gas from the combustion air, since oxygen and argon are co-emitted on the gas chromatograph. The magnitude of this last term is generally equal to 0.88 percent argon.

Carbon monoxide concentration is easily found by

$$C_{CO} = (C_{O_2} + 0.88) \frac{A_{CO}}{A_{O_2}} \quad (11)$$

which is simply ratioing the areas for the CO peak to that for the combined oxygen plus argon peak.

8.4 Correction of Measured Concentrations. Since much of the ducting routing flue gas through the air heater and subsequently into the stack is generally under negative pressure, there can be substantial leaks of ambient air into the ducting causing dilution of typically 10 to 15 percent, but sometimes more. In order to correct for this dilution so that simple calculation can then be made of the mass emission rate of pollutants based on the stoichiometry of the combustion, the concentrations found for SO_2 , H_2SO_4 , sulfate, and CO should be corrected by

$$C_{flue}^* = C_m \frac{20.95 - [O_2]_l}{20.95 - [O_2]_m} \quad (12)$$

where:

C_{flue}^* = concentration in flue gas with no leaks,

C_m = concentration as actually measured,

$[O_2]$ = oxygen level in furnace, vol percent, and

$[O_2]_m$ = measured oxygen, i.e., C_{O_2} from equation (10).

(Note: C_{SO_2} from (equation (12) should agree closely with the expected SO_2 level as calculated in Table 3.)

For H_2O , the correction is complicated slightly by the addition of water from the humidity of the ambient air that leaked into the duct. The correction is

$$[H_2O]_{flue}^* = \frac{[H_2O]_m(20.95 - [O_2]_l) - [H_2O]_{air}([O_2]_m - [O_2]_l)}{20.95 - [O_2]_m}$$

where:

$[H_2O]_{flue}^*$ = concentration of H_2O in the flue gas with no leaks, vol percent, and

$[H_2O]_{air}$ = concentration of H_2O in ambient air, vol percent.

Again, the water concentration as calculated above should agree with the expected level from Table 3.

The actual volume of furnace gas that was sampled, that is, the undiluted flue gas, can also be found by

$$V_{FW}^* = V_{FW} \frac{20.95 - [O_2]_m}{20.95 - [O_2]_l} \quad (14)$$

8.5 Total Suspended Particulate (TSP). The difference in weight of the NFA after use and its tare weight can be used to calculate TSP by

$$TSP = \frac{(W_{used} - W_{tare}) 10^6}{V_{FW}^*} \quad (15)$$

where:

TSP = total suspended particulate, mg/m^3 ,

W_{tare} = used weight of NFA, g,

V_{FW}^* = actual furnace gas volume, L (NTP).

Similarly, the amount of water insoluble fly ash in the flue gas can be calculated by

$$IP = \frac{(W_{wash} - W_{tare}) 10^6}{V_{FW}^*} \quad (16)$$

where:

IP = insoluble particulate, mg/m^3 , and

W_{wash} = weight of NFA after a water wash of filter, g.

8.6 NO_x Concentration. The NO_x concentration in the flue gas can be computed from the concentration found in the NO_x flask. First, the volume of the flue gas sample is given by

$$V_f = (V_f' - 0.01) \times \frac{P_f - P_i}{760} \times \frac{289}{(t + 273.2)} \quad (17)$$

where:

V_f = volume of sampled flue gas in NO_x flask, L,

V_f' = volume of flask, L,

P_f = final flask pressure after sampling, mm Hg,

P_i = initial flask pressure with acidified H_2O_2 , mm Hg, and

t = flask temperature during pressure measurement, °C.

The NO_x concentration in the sampled flue gas is given by

$$C_{NO_x} = \frac{C_{NO_3^-} \times 50 \times 23.7}{62 \times V_f} \quad (18)$$

where:

C_{NO_x} = NO_x concentration in sampled flue gas, vol ppm, and

$C_{NO_3^-}$ = nitrate concentration from ion chromatograph, $\mu g/mL$.

Taking into account the fact that the sampled flue gas had been dried and correcting for flue duct leaks, the actual NO_x concentration in the furnace gas is

$$C_{NO_x}^* = C_{NO_x} \times \frac{V_{FD}}{V_{FW}^*} \quad (19)$$

where:

$C_{NO_x}^*$ = NO_x concentration in furnace gas, vol ppm,

V_{FD} = volume of dry flue gas sampled (cf., equation (2)), L (NTP), and

V_{FW}^* = volume of wet flue gas with no ducting leaks (cf., equation (14)), L (NTP).

8.7 Calculated H_2SO_4 Dew Point Concentration. It is possible in some circumstances, especially when flue gas temperatures are too low, that the amount of H_2SO_4 in the flue gas can be close to the dew point concentration. Under such conditions, significant amounts of low temperature corrosion could occur. The dew point concentration can be calculated from (Reference 6).

$$\ln C_{H_2SO_4} = \frac{0.00002943 \ln(B.P. \times \frac{C_{H_2O}}{100} + \frac{1}{(t+273.2)}) - 0.002276}{0.0000858 [0.0723 \ln B.P. \times \frac{C_{H_2O}}{100} - 1]} - \ln \frac{B.P.}{10^6} \quad (20)$$

where:

$C_{H_2SO_4}$ = dew point concentration of H_2SO_4 , vol ppm,

C_{H_2O} = measured H_2O concentration (equation (7)), vol percent,

$B.P.$ = barometric pressure, mm Hg, and

t = flue gas temperature, $^{\circ}C$.

For example, at a flue gas temperature of $138^{\circ}C$ ($280^{\circ}F$), a water concentration of 12 percent, and a barometric pressure of 760 mm Hg, the dew point concentration is calculated to be 8.7 ppm H_2SO_4 . If a measured concentration comes close to the calculated dew point concentration, or exceeds it, then condensation of acid is almost certainly

occurring. That could lead to corrosion in the air heater and the ducting up to the stack.

8.8 Metals Emissions. The metals that were present in the flue gas fly ash can be expressed in terms of the portion that was present as water soluble sulfates and as insoluble oxides.

8.8.1 Water Soluble Metals

From the elemental analyses of the water soluble fraction of the filter washings, the concentration of soluble metals can be calculated by

$$C_{M-S} = \frac{50 M_s}{V_{FW}^*} \quad (21)$$

where:

C_{M-S}^* = concentration of soluble metals in flue gas, mg/m^3 ,

M_s = soluble metals concentration in 50 mL solution, $\mu g/mL$, and

V_{FW}^* = actual furnace gas volume, L (NTP).

8.8.2 Insoluble Metals

From the elemental analyses of the insoluble oxides remaining on the water washed filter, the concentration of insoluble metals can be calculated by

$$C_{M-I}^* = \frac{(IP \times V_{FW}^* + 90,000) M_I}{V_{FW}^*} 10^{-3} \quad (22)$$

where:

C_{M-I}^* = concentration of insoluble metals in flue gas, mg/m^3 ,

IP = insoluble particulate (from equation (16)), (that is, $\mu g/L$), and

M_I = insoluble metals concentration in ground filter, mg/g .

The term represented by 90,000 is the typical weight in μg of the 40 mm diameter filter.

Using this data on metal concentration in the flue gas, material balances can be performed on the inventory of metals distribution within the combustion unit (Reference 7).

8.9 Mass Emissions Calculations. Based on the stoichiometer equation for the combustion of the fuel, the total moles of furnace gas per mole of carbon can be calculated by an equation of the form of equation (2) in Table 3. This value, F , which will have a different equation for differences in fuel type and composition, can then be used to compute the volume of furnace gas produced per pound of fuel consumed:

$$F_v = 0.895 \times F \times [C] \quad (23)$$

where:

F_v = volume of furnace gas, $m^3/\#$ of fuel,

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F = moles of furnace gas/mole of carbon (cf., equation (2), Table 3), and

$[C]$ = wt fraction of carbon in the fuel.

Once F_v has been determined, it is a simple matter to compute mass emissions by multiplying F_v by the concentration in mass per unit volume of any of the previously calculated constituents. For example, the concentration of NO_x in mass per unit volume is

$$C_{\text{NO}_2}^{**} = C_{\text{NO}_x}^* \frac{MW_{\text{NO}_2}}{23.69} \quad (24)$$

where:

$C_{\text{NO}_2}^{**}$ = mass concentration expressed as mg of NO_2 per m^3 ,

$C_{\text{NO}_x}^*$ = NO_x concentration in furnace gas, vol ppm, and

MW_{NO_2} = molecular weight of NO_2 , 46 mg/mole.

The mass emission rate can then be calculated from

$$E_{\text{NO}_2} = 2.2 \times 10^{-6} F_v \times C_{\text{NO}_2}^{**} \quad (25)$$

where:

E_{NO_2} = pounds of NO_2 per pound of fuel.

Multiplying E_{NO_2} by the heating value of the fuel can convert E to mass emission per unit of heat input.

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FIG. 3

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HYDROGEN SULFIDE IN AIR

Physical and Chemical Analysis Branch

Analytical Method

Analyte:	Hydrogen Sulfide	Method No.:	P&CAM 126
Matrix:	Air	Range:	0.008 ppm – 50 ppm
Procedure:	Absorption – Methylene Blue – Spectrophotometric	Precision:	Unknown
Date Issued:	6/9/72	Classification:	C (Tentative)
Date Revised:	1/15/74		

1. Principle of the Method

- 1.1 Hydrogen sulfide is collected by aspirating a measured volume of air through an alkaline suspension of cadmium hydroxide (Reference 11.1). The sulfide is precipitated as cadmium sulfide to prevent air oxidation of the sulfide which occurs rapidly in an aqueous alkaline solution. STRactan 10[®] is added to the cadmium hydroxide slurry to minimize photo-decomposition of the precipitated cadmium sulfide (Reference 11.2). The collected sulfide is subsequently determined by spectrophotometric measurement of the methylene blue produced by the reaction of the sulfide with a strongly acid solution of N, N-dimethyl-p-phenylenediamine and ferric chloride (References 11.3, 11.4, 11.5). The analysis should be completed within 24-26 hours following collection of the sample.
- 1.2 Hydrogen sulfide may be present in the open atmosphere at concentrations of a few ppb or less. The reported odor detection threshold is in the 0.7-8.4 $\mu\text{g}/\text{m}^3$ (0.5-6.0 ppb) range (References 11.6, 11.7). Concentrations in excess of 140 $\mu\text{g}/\text{m}^3$ (100 ppb) are seldom encountered in the atmosphere.
- 1.3 Collection efficiency is variable below 10 $\mu\text{g}/\text{m}^3$ and is affected by the type of scrubber, the size of the gas bubbles and the contact time with the absorbing solution and the concentration of H_2S (References 11.8, 11.9, 11.10).

2. Range and Sensitivity

- 2.1 This method is intended to provide a measure of hydrogen sulfide in the range of 1.1-100 $\mu\text{g}/\text{m}^3$. For concentrations above 70 $\mu\text{g}/\text{m}^3$ the sampling period can be reduced or the liquid volume increased either before or after aspirating. (This

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method is also useful for the mg/m^3 range of source emissions. For example, 100 mL cadmium $(\text{OH})_2$ -STRactan 10[®] media in Greenberg-Smith impingers and 5 minute sampling periods have been used successfully for source sampling.) The minimum detectable amount of sulfide is $0.008 \mu\text{g/mL}$, which is equivalent to $0.2 \mu\text{g/m}^3$ in an air sample of 1 m^3 and using a final liquid volume of 25 mL. When sampling air at the maximum recommended rate of 1.5 L/minute for 2 hours, the minimum detectable sulfide concentration is $1.1 \mu\text{g/m}^3$ at 760 mm Hg and 25°C.

- 2.2 Excellent results have been obtained by using this method for air samples having a hydrogen sulfide content in the range 5-50 ppm.

3. Interferences

- 3.1 The methylene blue reaction is highly specific for sulfide at the low concentrations usually encountered in ambient air. Strong reducing agents (e.g., SO_2) inhibit color development. Even sulfide solutions containing several micrograms sulfide/mL show this effect and must be diluted to eliminate color inhibition. If sulfur dioxide is absorbed to give a sulfite concentration in excess of $10 \mu\text{g/mL}$, color formation is retarded. Up to $40 \mu\text{g/mL}$ of this interference, however, can be overcome by adding 2-6 drops (0.5 mL/drop) of ferric chloride instead of a single drop for color development, and extending the reaction time to 50 minutes.
- 3.2 Nitrogen dioxide gives a pale yellow color with the sulfide reagents at $0.5 \mu\text{g/mL}$ or more. No interference is encountered when 0.3 ppm NO_2 is aspirated through a midget impinger containing a slurry of cadmium hydroxide-cadmium sulfide-STRactan 10[®]. If H_2S and NO_2 are simultaneously aspirated through cadmium hydroxide-STRactan 10[®] slurry, lower H_2S results are obtained, probably because of gas phase oxidation of the H_2S prior to precipitation as CdS (Reference 11.10).
- 3.3 Ozone at 57 ppb reduced the recovery of sulfide previously precipitated as CdS by 15 per cent (Reference 11.10).
- 3.4 Sulfides in solution are oxidized by oxygen from the atmosphere unless inhibitors such as cadmium and STRactan 10[®] are present.
- 3.5 Substitution of other cation precipitants for the cadmium in the absorbent (i.e., zinc, mercury, etc.) will shift or eliminate the absorbance maximum of the solution upon addition of the acid-amine reagent.
- 3.6 Cadmium sulfide decomposes significantly when exposed to light unless protected by the addition of 1 per cent STRactan[®] to the absorbing solution prior to sampling (Reference 11.2).

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3.7 The choice of impinger used to trap H_2S with the $\text{Cd}(\text{OH})_2$ slurry is very important when measuring concentration in the range 5-50 ppm. Impingers or bubblers having fritted-end gas delivery tubes are a problem source if the sulfide in solution is oxidized by oxygen from the atmosphere to free sulfur. The sulfur collects on the fritted-glass membrane and may significantly change the flow rate of the air sample through the system. One way of avoiding this problem is to use a midjet impinger with standard, glass-tapered tips.

4. Precision and Accuracy

4.1 A relative standard deviation of 3.5 per cent and a recovery of 80 per cent has been established with hydrogen sulfide permeation tubes (Reference 11.2).

5. Advantages and Disadvantages of the Method

5.1 Effect of Light and Storage – Disadvantage

5.1.1 Hydrogen sulfide is readily volatilized from aqueous solution when the pH is below 7.0. Alkaline, aqueous sulfide solutions are very unstable because sulfide ion is rapidly oxidized by exposure to the air.

5.1.2 Cadmium sulfide is not appreciably oxidized even when aspirated with pure oxygen in the dark. However, exposure of an impinger containing cadmium sulfide to laboratory or to more intense light sources produces an immediate and variable photo-decomposition. Losses of 50-90 per cent of added sulfide have been routinely reported by a number of laboratories. Even though the addition of STRactan 10[®] to the absorbing solution controls the photo-decomposition (Reference 11.2), it is necessary to protect the impinger from light at all times. This is achieved by the use of low actinic glass impingers, paint on the exterior of the impingers, or an aluminum foil wrapping.

6. Apparatus

6.1 **Sampling Equipment.** The sampling unit for the impinger collection method consists of the following components:

6.1.1 A graduated 25 ml midjet impinger with a standard glass-tapered gas delivery tube containing the absorbing solution or reagent.

6.1.2 A pump suitable for delivering desired flow rates with a minimum capacity of 2 lpm through the impinger. The sampling pump is protected from splasher or water condensation by an adsorption tube loosely packed with a plug of glass wool and inserted between the exit arm of the impinger and the pump.

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6.1.3 An integrating volume meter such as a dry gas or wet test meter. The air meter must be capable of measuring the air flow within ± 2 per cent. A wet or dry gas meter, with contacts on the 1-foot³ or 10-l dial to record air volume, or a specially calibrated rotameter, is satisfactory. Instead of these, calibrated hypodermic needles may be used as critical orifices if the pump is capable of maintaining greater than 0.7 atmospheric pressure differential across the needle (Reference 11.11).

6.1.4 Thermometer.

6.1.5 Manometer.

6.1.6 Stopwatch.

6.2 Associated laboratory glassware.

6.3 Colorimeter with red filter or spectrophotometer at 670 nm.

6.4 Matched cells, 1 cm path length.

7. Reagents

All reagents must be ACS analytical reagent quality. Distilled water should conform to the ASTM Standards for Referee Reagent Water.

All reagents should be refrigerated when not in use.

7.1 **Amine-sulfuric Acid Stock Solution.** Add 50 ml concentrated sulfuric acid to 30 ml water and cool. Dissolve 12 g of N, N-dimethyl-p-phenylenediamine dihydrochloride (para-aminodimethylaniline) (redistilled if necessary) in the acid. Do not dilute. The stock solution may be stored indefinitely under refrigeration.

7.2 **Amine Test Solution.** Dilute 25 ml of the Stock Solution to 1 liter with 1:1 sulfuric acid.

7.3 **Ferric Chloride Solution.** Dissolve 100 g of ferric chloride, $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$, in water and dilute to 100 ml.

7.4 **Ammonium Phosphate Solution.** Dissolve 400 g of diammonium phosphate, $(\text{NH}_4)_2\text{HPO}_4$, in water and dilute to 1 liter.

7.5 **STRactan 10[®]**, (Arabinogalactan) Available from Stein-Hall and Company, Inc., 385 Madison Avenue, New York, New York.

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7.6 Absorbing Solution. Dissolve 4.3 g of $3\text{CdSO}_4 \cdot 8\text{H}_2\text{O}$, and 0.3 g sodium hydroxide in separate portions of water. Mix and add 10 g STRactan 10[®] and dilute to 1 liter. Shake the resultant suspension vigorously before removing each aliquot. The STRactan[®]-cadmium hydroxide mixture should be freshly prepared. The solution is only stable for 3 to 5 days.

7.7 H₂S Permeation Tube. Prepare or purchase a triple-walled or thick walled Teflon[®] permeation tube (References 11.10, 11.12, 11.13, 11.14, 11.15) which delivers hydrogen sulfide at a maximum rate of approximately 0.1 $\mu\text{g}/\text{minute}$ at 25°C. This loss rate will produce a standard atmosphere containing 50 $\mu\text{g}/\text{m}^3$ (36 ppb H₂S when the tube is swept with a 2 ℓ/minute air flow. Tubes having H₂S permeation rates in the range of 0.004-0.33 $\mu\text{g}/\text{minute}$ will produce standard air concentrations in the realistic range of 1-90 $\mu\text{g}/\text{m}^3$ H₂S with an air flow of 1.5 ℓ/min .

7.7.1 Concentrated, Standard Sulfide Solution

Transfer freshly boiled and cooled 0.1M NaOH to a liter volumetric flask. Flush with purified nitrogen to remove oxygen and adjust to volume. (Commercially available, compressed nitrogen contains trace quantities of oxygen in sufficient concentration to oxidize the small concentrations of sulfide contained in the standard and dilute standard sulfide standards. Trace quantities of oxygen should be removed by passing the stream of tank nitrogen through a Pyrex or quartz tube containing copper turnings heated to 400-450°C.) Immediately stopper the flask with a serum cap. Inject 300 ml of H₂S gas through the septum. Shake the flask. Withdraw measured volumes of standard solution with a 10 ml hypodermic syringe and fill the resulting void with an equal volume of nitrogen. Standardize with standard iodine and thiosulfate solution in an iodine flask under a nitrogen atmosphere to minimize air oxidation. The approximate concentration of the sulfide solution will be 440 μg sulfide/ml of solution. The exact concentration must be determined by iodine-thiosulfate standardization immediately prior to dilution.

For the most accurate results in the iodometric determination of sulfide in aqueous solution, the following general procedure is recommended:

1. Replace the oxygen from the flask by flushing with an inert gas such as carbon dioxide or nitrogen.

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2. Add an excess of standard iodine, acidify, and back titrate with standard thiosulfate and starch indicator (Reference 11.16).

7.7.2 Diluted Standard Sulfide Solution

Dilute 10 mL of the concentrated sulfide solution to 1 liter with freshly boiled, distilled water. Protect the boiled water under a nitrogen atmosphere while cooling. Transfer the deoxygenated water to a flask previously purged with nitrogen and immediately stopper the flask. This sulfide solution is unstable. Therefore, prepare this solution immediately prior to use. This test solution will contain approximately 4 µg sulfide/mL.

8. Procedure

- 8.1 Cleaning of Equipment. All glassware should be thoroughly cleaned; the following procedure is recommended:

- 8.1.1 Wash with a detergent and tap water solution followed by tap water and distilled water rinses.

- 8.1.2 Soak in 1:1 or concentrated nitric acid for 30 minutes and then follow with tap, distilled, and double distilled water rinses.

8.2 Collection and Shipping of Samples

- 8.2.1 Pipet 10 mL of the absorbing solution (Section 7.6) into the midjet impinger. The addition of 5 mL of 95 per cent ethanol to the absorbing solution just prior to aspiration controls foaming for 2 hours (induced by the presence of STRactan 10[®]). In addition, 1 or 2 Teflon demister discs may be slipped up over the impinger air inlet tube to a height approximately 1-2" from the top of the tube.

- 8.2.2 Connect the impinger (via the absorption tube) to the vacuum pump with a short piece of flexible tubing. The minimum amount of tubing necessary to make the joint between the prefilter and impinger should be used. The air being sampled should not be passed through any other tubing or other equipment before entering the impinger.

- 8.2.3 Turn on pump to begin sample collection. Care should be taken to measure the flow rate, time and/or volume as accurately as possible. The sample should be taken at a flow rate of 1.5 Lpm.

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- 8.2.4 After sampling, the impinger stem can be removed and cleaned. Tap the stem gently against the inside wall of the impinger bottle to recover as much of the sampling solution as possible. Wash the stem with a small amount (1-2 mL) of unused absorbing solution and add the wash to the impinger. Then the impinger is sealed with a hard, non-reactive stopper (preferably Teflon). Do not seal with rubber. The stoppers on the impingers should be tightly sealed to prevent leakage during shipping. If it is preferred to ship the impingers with the stems in, the outlets of the stem should be sealed with Parafilm or other non-rubber covers, and the ground glass joints should be sealed (i.e., taped) to secure the top tightly.
- 8.2.5 Care should be taken to minimize spillage or loss by evaporation at all times. Refrigerate samples if analysis cannot be done within a day.
- 8.2.6 Whenever possible, hand delivery of the samples is recommended. Otherwise, special impinger shipping cases designed by NIOSH should be used to ship the samples.
- 8.2.7 A "blank" impinger should be handled as the other samples (fill, seal and transport) except that no air is sampled through this impinger.

8.3 Analysis

- 8.3.1 Add 1.5 mL of the amine-test solution to the midget impinger through the air inlet tube and mix.
- 8.3.2 Add 1 drop of ferric chloride solution and mix. (*Note:* See Section 3.1 if SO_2 exceeds $10 \mu\text{g/mL}$ in the absorbing media.)
- 8.3.3 Transfer the solution to a 25 mL volumetric flask. Discharge the color due to the ferric ion by adding 1 drop ammonium phosphate solution. If the yellow color is not destroyed by 1 drop ammonium phosphate solution, continue dropwise addition until solution is decolorized. Make up to volume with distilled water and allow to stand for 30 minutes.
- 8.3.4 Prepare a zero reference solution in the same manner using a 10 mL volume of absorbing solution, through which no air has been aspirated.
- 8.3.5 Measure the absorbance of the color at 670 nm in a spectrophotometer or colorimeter set at 100 per cent transmission against the zero reference.

9. Calibration and Standards

9.1 Aqueous Sulfide

- 9.1.1 Place 10 mL of the absorbing solution in each of a series of 25 mL volumetric flasks and add the diluted standard sulfide solution, equivalent to 1, 2, 3, 4, and 5 μg of hydrogen sulfide to the different flasks.
- 9.1.2 Add 1.5 mL of amine-acid test solution to each flask and mix.
- 9.1.3 Add 1 drop of ferric chloride solution to each flask. Mix, make up to volume and allow to stand for 30 minutes.
- 9.1.4 Determine the absorbance in a spectrophotometer at 670 nm, against the sulfide-free reference solution.
- 9.1.5 Prepare a standard curve of absorbance vs. $\mu\text{g H}_2\text{S/mL}$.

9.2 **Gaseous Sulfide.** Commercially available permeation tubes containing liquefied hydrogen sulfide may be used to prepare calibration curves for use at the upper range of atmospheric concentration. Triple-walled tubes, drilled rod and micro bottles which deliver hydrogen sulfide within a minimum range of 0.1-1.2 $\mu\text{g/minute}$ at 25°C have been prepared by Thomas (Reference 11.10); O'Keeffe (References 11.12, 11.13); Scaringelli (References 11.14, 11.15). Preferably the tubes should deliver hydrogen sulfide within a loss rate range of 0.003-0.28 $\mu\text{g/minute}$ to provide realistic concentrations of H_2S (1.5-140 $\mu\text{g/m}^3$, 1.1-100 ppb) without having to resort to a dilution system to prepare the concentration range required for determining the collection efficiency of midget impingers. Analyses of these known concentrations give calibration curves which simulate all of the operational conditions performed during the sampling and chemical procedure. This calibration curve includes the important correction for collection efficiency at various concentrations of hydrogen sulfide.

- 9.2.1 Prepare or obtain a Teflon[®] permeation tube that emits hydrogen sulfide at a rate of 0.1-0.2 $\mu\text{g/minute}$ (0.07-0.14 $\mu\text{L/minute}$ at standard conditions of 25°C and 1 atmosphere). A permeation tube with an effective length of 2-3 cm and a wall thickness of 0.318 cm will yield the desired permeation rate if held at a constant temperature of 25°C $\pm$ 0.1°C. Permeation tubes containing hydrogen sulfide are calibrated under a stream of dry nitrogen to prevent the precipitation of sulfur in the walls of the tube.
- 9.2.2 To prepare standard concentrations of hydrogen sulfide, assemble the apparatus consisting of a water-cooled condenser, constant temperature bath maintained at 25°C $\pm$ 0.1°C cylinders containing pure dry nitrogen

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and pure dry air with appropriate pressure regulators, needle valves and flow meters for the nitrogen and dry air, diluent-streams. The diluent gases are brought to temperature by passage through a 2-meter-long copper coil immersed in the water bath. Insert a calibrated permeation tube into the central tube of the condenser, maintained at the selected constant temperature by circulating water from the constant-temperature bath, and pass a stream of nitrogen over the tube at a fixed rate of approximately 50 mL/minute. Dilute this gas stream to obtain the desired concentration by varying the flow rate of the clean, dry air. This flow rate can normally be varied from 0.2-15 L/minute. The flow rate of the sampling system determines the lower limit for the flow rate of the diluent gases. The flow rate of the nitrogen and the diluent air must be measured to an accuracy of 1-2 per cent. With a tube permeating hydrogen sulfide at a rate of 0.1 $\mu\text{L}/\text{minute}$, the range of concentration of hydrogen sulfide will be between 6-400 $\mu\text{g}/\text{m}^3$ (4-290 ppb), a generally satisfactory range for ambient air conditions. When higher concentrations are desired, calibrate and use longer permeation tubes.

9.2.3 Procedure for Preparing Simulated Calibration Curves

Obviously one can prepare a multitude of curves by selecting different combinations of sampling rate and sampling time. The following description represents a typical procedure for ambient air sampling of short duration, with a brief mention of a modification for 24 hour sampling.

1. The system is designed to provide an accurate measure of hydrogen sulfide in the 1.4-84 $\mu\text{g}/\text{m}^3$ (1-60 ppb) range. It can be easily modified to meet special needs.
2. The dynamic range of the colorimetric procedure fixes the total volume of the sample at 186 L; then, to obtain linearity between the absorbance of the solution and the concentration of hydrogen sulfide in ppm, select a constant sampling time. This fixing of the sampling time is desirable also from a practical standpoint: In this case, select a sampling time of 120 minutes. Then to obtain a 186 L sample of air requires a flow rate of 1.55 L/minute. The concentration of standard H_2S in air is computed as follows:

$$C = \frac{P_r \times M}{(R + r)}$$

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where:

- C = Concentration of H_2S in ppm
- P_r = Permeation rate in $\mu g/\text{minute}$
- M = Reciprocal of vapor density, $0.719 \mu\ell/\mu g$
- R = Flow rate of diluent air, liter/minute
- r = Flow rate of diluent nitrogen, liter/minute

3. The data for a typical calibration curve are listed in Table 1.

TABLE 1
TYPICAL CALIBRATION DATA

<u>Concentrations</u> <u>H_2S, ppb</u>	<u>Amount of</u> <u>H_2S in</u> <u>$\mu\ell/186$ liters</u>	<u>Absorbance</u> <u>of Sample</u>
1	.144	.010
5	.795	.056
10	1.44	.102
20	2.88	.205
30	4.32	.307
40	5.76	.410
50	7.95	.512
60	8.64	.615

4. A plot of the concentration of hydrogen sulfide in ppm (x – axis) against absorbance of the final solution (y – axis) will yield a straight line, the reciprocal of the slope of which is the factor for conversion of absorbance to ppm. This factor includes the correction for collection efficiency. Any deviation from the linearity at the lower concentration range indicates a change in collection efficiency of the sampling system. If the range of interest is below the dynamic range of the method the total volume of air collected should be increased to obtain sufficient color within the dynamic range of the colorimetric procedure. Also, once the calibration factor has been established under simulated conditions the conditions can be modified so that the concentration of H_2S is a simple multiple of the absorbance of the colored solution.

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5. For long-term sampling of 24-hour duration, the conditions can be fixed to collect 1200 l of sample in a larger volume of STRactan 10[®]-cadmium hydroxide. For example, for 24 hours at 0.83 l/min, approximately 1200 l of air are scrubbed. An aliquot representing 0.1 of the entire amount of sample is taken for the analysis.
6. The remainder of the analytical procedure is the same as described in the previous paragraph.

- 9.2.4 The permeation tubes must be stored in a wide-mouth glass bottle containing silica gel and solid sodium hydroxide to remove moisture and hydrogen sulfide. The storage bottle is immersed to two-thirds its depth in a constant temperature water bath in which the water is controlled at 25°C ±0.1°C.

Periodically, (every 2 weeks or less) the permeation tubes are removed and rapidly weighed on a semimicro balance (sensitivity ±0.01 mg) and then returned to the storage bottle. The weight loss is recorded. The tubes are ready for use when the rate of weight loss becomes constant (within ±2 per cent).

10. Calculations

10.1 Gaseous Sulfide

- 10.1.1 Determine the sample volume in liters from the gas meter or flow meter readings and time of sampling. Adjust volume to 760 mm mercury and 25°C (V_s).

- 10.1.2 Determine the concentration of H_2S in $\mu g/m^3$.

$$\mu g H_2S/m^3 = \mu g H_2S/V_s \times 10^{-3}$$

where:

$$\begin{array}{ll} \mu g H_2S & = \text{micrograms hydrogen sulfide determined} \\ 10^{-3} & = \text{conversion factor, } m^3/l \end{array}$$

10.2 Gaseous Sulfide from Aqueous Sulfide

- 10.2.1 Determine the sample volume (V) in liters from the gas meter or flow meter readings and time of sampling. Adjust volume to 760 mm mercury and 25°C (V_s), using the correction formula:

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$$V_s = V \times \frac{P}{760} \times \frac{298}{(T + 273)}$$

where:

- V_s = Volume of air in liters at standard conditions
- V = Volume of air sampled in liters
- P = Barometric pressure in mm Hg
- T = Temperature of sample air in °C

10.2.2 Using the Beers-Law Standard curve of absorbance vs. $\mu\text{g S}^{=}$ ion, determine $\mu\text{g S}^{=}$ ion in the sampling impinger corresponding to its absorbance reading at 670 nm.

10.2.3 Calculate the concentration of H_2S in the aspirated volume of air using the formula:

$$\text{ppm, H}_2\text{S} = \frac{\mu\text{g S}^{=} \times 24.45}{V_s \times \text{MW}}$$

where:

- $\mu\text{g S}^{=}$ = micrograms sulfide ion (Section 10.2.2)
- 24.45 = molar volume of an ideal gas at 25°C and 760 mm Hg
- MW = mass of sulfide ion, 32.06

11. References

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