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Metode vzorčenja in analize dimnih plinov

Methods for sampling and analysis of fire effluents

iTeh STANDARD PREVIEW
Méthodes d'échantillonnage et d'analyse des effluents du feu
(standards.iteh.ai)

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**Methods for sampling and analysis of
fire effluents**

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Case postale 56 • CH-1211 Geneva 20
Tel. + 41 22 749 01 11
Fax + 41 22 749 09 47
E-mail copyright@iso.org
Web www.iso.org

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ISO 19701:2013(E)**Foreword**

ISO (the International Organization for Standardization) is a worldwide federation of national standards bodies (ISO member bodies). The work of preparing International Standards is normally carried out through ISO technical committees. Each member body interested in a subject for which a technical committee has been established has the right to be represented on that committee. International organizations, governmental and non-governmental, in liaison with ISO, also take part in the work. ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization.

International Standards are drafted in accordance with the rules given in the ISO/IEC Directives, Part 2.

The main task of technical committees is to prepare International Standards. Draft International Standards adopted by the technical committees are circulated to the member bodies for voting. Publication as an International Standard requires approval by at least 75 % of the member bodies casting a vote.

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. ISO shall not be held responsible for identifying any or all such patent rights.

ISO 19701 was prepared by Technical Committee ISO/TC 92, *Fire safety*, Subcommittee SC 3, *Fire threat to people and environment*.

This second edition cancels and replaces the first edition (ISO 19701:2005).

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Introduction

The analysis of fire gases for use in toxic hazard and life threat assessment and other purposes (e.g. impact on the environment) is a great challenge for the analyst. Fire atmospheres are by nature hostile environments. Temperatures in excess of 1 000 °C are common, the gas phase can contain many corrosive, toxic, irritant or combustible species together with relatively large quantities of condensable water.

These properties are largely incompatible with most instrumental analytical methods where a “clean” sample is required. This poses many problems both for the qualification and quantification of the chemical species and particulates in fire atmospheres. In presenting a sample to the measuring instrument that it will tolerate, it can be necessary to filter particulates and remove other species. Losses in the sampling train must therefore be quantifiable and taken into account in the final analysis.

Techniques also exist for measuring chemical species *in situ*; this will be the subject of a future document.

The methods described in [Clause 5](#) have been used successfully by a number of laboratories. Studies of repeatability and reproducibility of many of the methods covered in this International Standard have been taken from AFNOR NF X70-100-1[1] and AFAP-3.[2]

For methods that involve a commercial instrument, uncertainty in the measured values may be estimated from the manufacturer’s data and other information, e.g. allowance for losses in the sampling process. For other methods, uncertainty in the measured values can occur through a variety of reasons, such as sensitivity to the strength of reagents or the visibility of a colourimetric end point. In these cases, it is assumed that best practice by qualified personnel is applied.

This International Standard is structured as follows.

- [Clause 1](#) describes the scope of this standard
- [Clause 4](#) describes methods of sampling
- [Clause 5](#) describes analytical methods for gases in fire atmospheres:
- [Annex A](#) provides information on techniques that were found not suitable with fire effluents.
- [Annex B](#) briefly describes the use of aspirated chemical colour-change tubes.
- [Annex C](#) is a summary of the main instrumental methods available for fire gas analysis, expanding the information provided under the clauses for each individual chemical species.
- [Annex D](#) presents a method for continuous measurement of HF concentration using ion selective electrode.

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Methods for sampling and analysis of fire effluents

SAFETY PRECAUTIONS — Due consideration must be given to the fact that both the fire gases for analysis and many of the reagents used for their analysis can be toxic and/or present serious health hazards. It is assumed throughout that the procedures described in this document will be carried out by suitably qualified professional personnel, adequately trained in the hazards and risks associated with such analyses and aware of any safety regulations that may be in force. Consideration must also be given to the safe and ecologically acceptable disposal of all chemicals used for analyses. This can require extensive treatment prior to release of the waste into the environment. Again, it is assumed in this document that the personnel responsible for the safe disposal of such reagents are suitably qualified and trained in these techniques and are aware of the regulations which may be in force.

1 Scope

This International Standard presents a range of sampling and chemical analytical methods suitable for the analysis of individual chemical species in fire atmospheres. The procedures relate to the analysis of samples extracted from an apparatus or effluent flow from a fire test rig or physical fire test model and are not concerned with the specific nature of the fire test.

This International Standard doesn't cover aerosols (detailed in Reference [3]) and FTIR technique (detailed in Reference [4]). The gases of environmental interest, such as PAH, dioxins, furans, endocrinal disturbers, will be developed in a future document by ISO TC92/SC3.

2 Normative references

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The following documents, in whole or in part, are normatively referenced in this document and are indispensable for its application. For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments) applies.

ISO 13943, *Fire safety — Vocabulary*

3 Terms and definitions

For the purposes of this document, the terms and definitions given in ISO 13943 apply.

4 Methods of sampling

4.1 General considerations

Sampling is perhaps the most critical part of the procedures for analysis of gases in fire effluents. Whereas sampling and analysis are commonly in use for many gaseous species in other fields, sampling from fire atmospheres presents unusual and difficult problems.

The sample presented to the analyser shall be as representative as possible of the test atmosphere, without having been changed by the sampling system. The sampling procedure should influence the test atmosphere as little as possible (e.g. by depletion of the test volume). The sampling procedure should be as uncomplicated as possible, while incorporating all necessary features detailed in this International Standard. The sampling procedure shall be capable of operating with minimal blockage in the sampling lines, melting or other disruption of probes, and without allowing condensation of the species for analysis.

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IMPORTANT — It is important to appreciate that the overall accuracy of the analysis of fire effluent species is significantly dependent on the sampling procedures adopted, in particular the quantification of losses in probes, sampling lines, and filtering systems.

4.2 Concentration and volume fraction

The concentration of fire effluent or of a toxic gas is its mass divided by the volume in which it is contained. For a fire effluent the typical units are $\text{g}\cdot\text{m}^{-3}$. However, for a toxic gas, concentration is usually expressed as a volume fraction at $T = 298 \text{ K}$ and $P = 1 \text{ atm}$, and is expressed in terms of $\mu\text{L}\cdot\text{L}^{-1}$ (equivalent to $\text{cm}^3/\text{m}^3 = 10^{-6}$).

NOTE 1 The concentration of a gas at a temperature, T , and a pressure, P can be calculated from its volume fraction (assuming ideal gas behaviour) by multiplying the volume fraction by the density of the gas at that temperature and pressure.

NOTE 2 Volume fractions of toxic gases used to be expressed in terms of “ppm by volume” but “ppm” is a deprecated term and therefore “ $\mu\text{L}\cdot\text{L}^{-1}$ ” is now used.

4.3 Special considerations

There are many factors that have a direct influence on the specific type of sampling methodology selected to ensure that a suitable sample is presented to the analyser. For example, consideration shall include the range of concentrations anticipated, the limits of detection, reactivity of the species of interest, presence of interferences, and peak and average concentration values. Sampling of the extremely complex atmosphere produced during combustion requires a very thorough evaluation and assessment of all potential factors that might affect optimum conditions for sample collection and analysis.

The large number of different products frequently encountered in fire effluents often requires the use of a variety of sampling procedures and approaches to ensure accurate identification and quantification of combustion products. The selected sampling procedure also depends on the instrumentation and analytical procedures available for the specific species being measured.

Sampling may involve either continuous, online analysis (e.g. non-dispersive infrared) or non-continuous batch sampling (e.g. evacuated flask or bubbler samples). Batch-type sampling can be further subdivided into two categories:

- a) “Instantaneous”, or “grab”;
- b) Average, or integrated.

Although there is no sharp distinction between categories a) and b), it is generally understood that grab samples relate to samples taken over a short time period (i.e. usually less than 1 min), whereas integrated samples are usually taken over a longer time period (i.e. a substantial portion of the total test period).

In some cases, continuous or semi-continuous online or frequent instantaneous sampling can be well suited for following the rapidly changing combustion environment and will provide a representative concentration profile. Frequently however, the minimum detectable limit of the species of interest requires larger sample volumes than can be taken with these techniques. If this analytical limitation exists, it is necessary to carry out the sampling over a longer period. While using longer sampling periods permits the analysis of lower concentrations, this approach has some limitations. For example, these types of samples permit a determination only of the integrated average concentration obtained over the sampling period and do not discern any abrupt change in the evolution of the species of interest. However, abrupt concentration changes can also be missed with instantaneously obtained samples, if samples are not taken frequently enough.

When batch-sampling procedures are used, it is essential to specify sampling frequency, the starting time of each sample and the total sampling time. This information is essential in order to ensure proper evaluation of the data in conjunction with other fire properties that are being monitored (e.g. heat release, temperatures, mass loss, smoke evolution, flame spread).

Test fires can be classified as “small” (laboratory or “bench” size), “intermediate” or “large” (usually full-scale). The sampled gases can be hot or near room temperature. It is generally necessary to extract the gases from the test atmosphere through suitable tubing using a suction pump. Stainless steel tubing, as short as possible, is often used. In the case of the production of hot gases, the sampling line shall be heated to at least 100 °C. Several analytical methods require a dry, particulate-free sample. Glass wool may be used (in most cases) as a particulate filter, with another trap of a drying agent (e.g. calcium sulfate or calcium chloride) for removing moisture. The traps should be located just before the analyser and after any heated sections of sampling tubes. Simple cold traps are often insufficient to remove the quantity of moisture present in fire effluents; however, they can be useful in conjunction with other filters and traps. The individual sampling and analytical system being used dictates flow requirements and the necessity for moisture removal. Precautions should be taken to minimize the volume of the filtering systems to reduce sampling time.

With the exception of hydrogen fluoride (HF), acid gases shall be sampled using glass, epoxy-lined or PTFE tubes to minimize losses due to reactivity and condensation on the tube surfaces. For hydrogen fluoride, tubes lined with PTFE shall be used (glass and glass-lined tubes are unsuitable). For species that are relatively reactive and prone to losses, sampling lines shall be as short as possible, and shall be heated to a sufficient temperature to avoid condensation. Hydrogen chloride (HCl) and hydrogen bromide (HBr) can be adsorbed onto soot particles as well as gas sampling lines (including PTFE lines).

For organic materials (e.g. acrolein), unlined stainless steel tubing is suitable but the sampling lines shall be heated to avoid condensation. Particulate traps, although usually necessary, can be avoided in some cases and instrument requirements should be checked in this regard.

The location and size of sampling probes is influenced by the size of the test apparatus and the requirements of the analytical system. The positioning of sampling probes in specific apparatus, however, is beyond the scope of this International Standard. In general, the possibility of the stratification of gases in chambers without good mixing shall be considered and sampling too near the wall of a test chamber should be avoided.

Calibration of the ~~entire sampling and analysis system, rather than just the analysis system,~~ is recommended in order to ensure that any losses in the sample route can be allowed for. All calibrations should, therefore, take into account such factors as gas leakage (both into and out of the sampling lines) and the adsorption of gases onto probes, sampling lines, filters and other components. Calibration gases are often obtainable in cylinders; however, it is advisable that the concentration stated by the supplier be verified by an independent analysis. This is especially true of reactive gases such as HCl and HF, which can decay over relatively short time periods even in a closed cylinder. The calibration gas shall be introduced at the sampling probe and allowed to travel the same course as a test gas, through filters and traps if present, to the analyser or sampling medium.

4.4 Sampling using gas-solution absorbers

Absorption of gases in solution by the use of gas-washing bottles, bubblers, impingers, etc. all rely on the same principle. The test atmosphere is drawn or pushed through the absorbing media at a measured rate for a specified period of time. At the end of the sampling period, the solution is analysed for the species of interest (e.g. the chloride ion for absorption of hydrogen chloride gas in water). Assuming 100 % efficiency

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(see discussion below), it is possible to calculate the concentration of the species in the gas phase, as measured in the solution. A typical equation for calculating concentrations is presented in Formula (1):

$$\rho_G = \frac{\rho_S \times V \times (m_G / m_S)}{q \times t} \quad (1)$$

where

ρ_G is the gas concentration;

ρ_S is the solution concentration;

V is the volume, expressed in litres, of solution;

m_G/m_S is the ratio of atomic or molecular weights for the gaseous species, G , and solution species, S , if different, e.g. HCl/Cl);

q is the rate of gas flow, expressed in litres per minute, through the impinger;

t is the time, in minutes of gas flow.

The volume fraction of the gas, X_G , can be calculated by dividing the concentration by the density d , of the gas at the ambient temperature and pressure. This density can be found, assuming ideal gas behaviour, as follows:

$$d = \frac{M_G}{H} \quad (2)$$

$$X_G = \rho_G \frac{H}{M_G} \quad (3)$$

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where

M_G is the molar mass of the gaseous species, G ;

H is the gaseous volume occupied by 1 mol of an ideal gas at the relevant ambient pressure (P) and temperature (T). [$H = 8,314 \text{ J.K}^{-1}.\text{mol}^{-1} \times 1 \text{ mol} \times T/P$]

EXAMPLE Suppose that the measured solution concentration of chloride ion (Cl^-) was $0,006 \text{ g} \times \text{dm}^{-3}$ in $0,025 \text{ dm}^3$ of solution, the ambient thermodynamic temperature was $293,15 \text{ K}$, the ambient pressure was 1 bar ($10^5 \text{ Pa} = 10^5 \text{ J} \times \text{m}^{-3}$), and the flow rate of gas was $0,25 \text{ dm}^3/\text{min}$ for 2 min . Then the gas concentration of hydrogen chloride is given by:

$$\rho_G = [0,006 \text{ g}.\text{dm}^{-3} \times 0,025 \text{ dm}^3 \times (36,461 / 35,453)] / [0,25 \text{ dm}^3/\text{min} \times 2 \text{ min}]$$

$$\rho_G = [0,00015 \text{ g} \times 1,028] / 0,50 \text{ dm}^3$$

$$\rho_G = 0,0003084 \text{ g}.\text{dm}^{-3} = 308,4 \text{ mg}.\text{m}^{-3}$$

The volume fraction of hydrogen chloride is given by:

$$X_G = 308,4 \text{ mg}.\text{m}^{-3} \times (8,314 \text{ J.K}^{-1}.\text{mol}^{-1} \times 293,15 \text{ K} / 10^5 \text{ J}.\text{m}^{-3}) / 36,461 \text{ g}.\text{mol}^{-1}$$

$$X_G = 308,4 \text{ mg}.\text{m}^{-3} \times 0,02437 \text{ m}^3.\text{mol}^{-1} / 36,461 \text{ g}.\text{mol}^{-1}$$

$$X_G = 0,0002061 = 206,1 \mu\text{L}/\text{L}$$

The volume of the absorber solution and the total flow of gas directly affect the ratio of the gas and solution concentrations. For a given gas concentration, the smaller the solution volume and/or the larger the gas volume sampled, the higher the solution concentration. The choice of sampling conditions

is dictated by the requirements of the analytical technique, including the volume and sampling rate tolerated, expected concentration of gas in the test atmosphere, necessity for frequent sampling, etc.

The efficiency of absorption of a gas in liquid is affected by the following:

- a) Solubility of the gas in the solution;
- b) Physical characteristics of the absorber;
- c) Ratio of gas flow rate to solution volume.

Generally, absorption efficiency is estimated empirically by allowing the flow of a known concentration of the gas of interest through a series of impingers and measuring the “break-through” from the first impinger (i.e. whatever is collected in the other traps). Another check on the efficiency of a given flow/impinger system is to conduct a series of experiments with a known concentration of gas, using different impingers and various flow rates. In practice, however, the choice of apparatus is limited, and gas flow rates and trapping solution volumes are based on Formula (1), taking into account the known characteristics of the analysis methods.

There are basically four types of gas-solution absorbers: simple gas-washing bottles (including midget impingers), spiral or helical absorbers, packed glass-bead columns and fritted bubblers. The gas-washing bottles, or impingers, function by drawing the gas through a tube (usually with a constricted opening), which is immersed in the trapping liquid/solution. This type is most suitable for highly soluble gases because contact time between solution and gas is short and bubble size is relatively large. For less soluble species, the other absorbers offer longer contact time and/or smaller bubble size (which increases relative surface contact). The spiral or helical absorbers are built in specialized shapes to allow a long contact time. The flow rate in these bubblers is limited because of the possibility of trapping solution over-flow with high flow rates. Packed glass-bead columns allow increased gas/liquid contact by dispersing the bubbles through a bed of glass beads. Flow rates can be higher than for the spiral absorbers.

The fritted bubblers contain a sintered or fritted disc on the gas inlet tube to disperse the gas into fine bubbles (the size of the bubbles is dependent on the porosity of the frit). It is necessary to exercise caution in using such bubblers so that frothing does not occur and so that the coalescence of the fine bubbles does not defeat the purpose of the frit. Also, it is necessary to filter smoky atmospheres (containing particulates or liquid aerosols) before drawing them through a fritted bubbler in order to prevent clogging of the frit (which occurs very easily). Such clogging can also occur from the build-up of wax-like deposits. Certain gas species (e.g. HCl) can be absorbed onto a filter, especially if particulates have also been trapped on the filter.

Note that very soluble gases, such as HCl and HF, can cause water to be sucked back along the sampling tube. With these gases, it is often necessary to include an empty bubbler to act as a liquid trap.

4.5 Sampling using solid sorption tubes

Solid sorption tubes are an alternative method to gas-solution absorbers for sampling certain gases from fire effluents. Following sampling, the species of interest is desorbed in water and its analysis can then be performed in a way similar to that for aqueous solution absorbers.

The advantages of solid sorption tubes over solution absorbers are

- a) Ease of handling,
- b) Compactness,
- c) High absorption efficiency,
- d) Ability to be located directly at the point of sampling.

This latter advantage can have dramatic consequences in the measurement of HF, HCl and HBr in fire effluents because these species are easily lost to the inside surfaces of sampling lines. With solid sorption tubes (except in areas of extreme heat), a sampling line is not necessary before the sorption tube itself. All

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associated hardware (e.g. valves, flow meters and pumps) can be located behind the tubes, even far from the sampling point. This ensures that the sample is as representative as possible of the fire atmosphere.

Much experience has been gained through using solid sorption tubes, for example in the field of atmospheric sampling and for staff exposure monitoring in the workplace. Similar tubes have been re-examined for potential use in sampling fire effluents. Two studies [5],[6] were carried out using solid sorbents to measure certain gases in real building fires. These tubes were located in portable sampling boxes carried by the firemen who were actually fighting the fire. Tubes of similar design, containing activated charcoal, have been used to sample HF [7] and HCN.[8] Tubes containing flake sodium hydroxide for the absorption of acid gases have also been described.[9] A procedure for successive (e.g. every 3 min or 5 min) sampling with tubes at one location without removing or replacing tubes has been described for sampling gases in full-scale fires.[8]

Calculation of the original gas concentration (e.g. HCl) from the representative species recovered in the desorbent solution (e.g. Cl⁻) is the same as that described for solution absorbers, except that the solution volume is the volume of desorbent liquid. In practice, a small aliquot, rather than the entire quantity, of the desorbent solution is often used for the analysis so it is necessary to take this factor into account.

The same considerations that apply to solution absorbers, with respect to inefficient absorption, breakthrough and the relationship of volume sampled to gas and solution concentration, also apply to the use of solid sorbents. Instead of bubble size, it is the particulate size of the absorbent that is important (large particles offer less surface area per unit volume and more opportunity for channelling, smaller particles can cause the tube to plug when sampling moist gas). The tubes should be small enough (typically 100 mm long, 6 mm OD) such that two tubes can easily be placed in series to allow for the possibility of "breakthrough" from the first tube.

Solid sorption tubes are subject to plugging due to soot collection. This can be recognized during sampling by a decrease in sample flow rate. The same flow rate should be maintained over the duration of sampling using a constant flow device; otherwise, an error is introduced in the calculation of gas concentration. A glass wool plug loosely packed into the inlet of the tube reduces the tendency to blocking from soot.

Thermal desorption of the adsorbed sample is also possible; the sample tube is heated in an inert gas stream thus driving off the sample without the need for a liquid solution stage.

4.6 Sampling for spectrometric or spectrophotometric analysis

The uses of spectrometric analysis [direct mass spectrometry (MS)] and spectrophotometric analysis [both non-dispersive infrared (NDIR) and Fourier transform infrared (FTIR)] have become quite widespread in recent years. FTIR techniques in particular are becoming more prominent.[9], [10], [11] The continuous measurement by means of NDIR analysis (e.g. for CO and CO₂) is now so common that several different companies manufacture commercial instruments designed for this purpose.

For two of the methods (direct MS and FTIR), it is important that the fire effluents be free from particles before they are introduced into the analyser. The filter used, which is often placed at the junction of the sampling line and the test chamber, shall be inert so it does not react with any of the gases of interest. A stainless steel filter unit containing a glass-fibre filter (e.g. Whatman multigrade GMF150¹) micro-filter, 1 µm, 47 mm in diameter) has been found suitable. The sample line and the filter (and for FTIR also the absorption cell) are heated to a temperature above 120 °C (120 °C to 150 °C has been found to be suitable), in order to prevent liquid water from forming, to prevent water-soluble gases (e.g. HCN and the acid gases) from dissolving and other gases from condensing.

When a filter is used, it is necessary to check the extent to which the species of interest have been retained by the filter. If retention occurs, it is necessary to correct the measured concentrations. The amount of retained material is dependent principally on the type and capacity of the filter used, the nature of the species and the volume of gas passing through the filter.

1) The Whatman GMF150 filter is an example of a suitable product available commercially. This information is given for the convenience of users of ISO 19701 and does not constitute an endorsement by ISO of this product.

4.7 Sampling using gas bags

Sampling with gas bags can be used for most analytical methods. The test atmosphere is pumped, or allowed to flow under pressure, into a gas bag at a measured constant rate for a measured time period, thus obtaining a known volume of sample in the bag. It is necessary to filter the fire effluents before passing into the bag; simple in-line glass wool filters for particulates, and calcium chloride filters for moisture, have been found effective. However, a calcium chloride absorbent removes water vapour and water-soluble gases. At the end of the sampling period, the bag may be stored before it is connected to the analyser; but it is important to appreciate that storage times in bags should be kept to a minimum, preferably less than 1 h. Gases such as HF and HCl can dissolve in condensed/trapped water and this reduces the concentration presented to the analyser.

Bags shall be gas-tight and inert and those with a lining of polyvinylfluoride (PVF) are recommended.

[Table 1](#) summarizes the analytical methods and types of sample required for each method described in this International Standard.

Table 1 — Type of sampling for the analytical methods described

Gas	Analytical Method	Type of sample for analysis
Carbon monoxide (CO)	NDIR	gas
Carbon dioxide (CO ₂)	NDIR	gas
Oxygen (O ₂)	Paramagnetism	gas
Hydrogen cyanide(HCN)	Colourimetry (Chloramine T) Colourimetry (picric acid) HPIC	solution solution solution
Hydrogen chloride (HCl) Hydrogen bromide (HBr)	ISE HPIC titrimetry	solution solution solution
Hydrogen fluoride (HF)	ISE HPIC Online ISE	solution solution solution
Nitrogen oxides (NO _x)	Chemiluminescence	gas
Nitrogen dioxide (NO ₂)	HPIC	solution
Nitrogen monoxide (NO)	Gfx-IR	gas
Acrolein (2-propenal)	Colourimetry HPLC GC-MS	solution solution gas
Formaldehyde (Methanal)	Colourimetry HPLC	solution solution
Acetaldehyde (Ethanal)	HPLC GC-MS	solution solution
Total aldehydes	Colourimetry	solution
Sulfur dioxide (SO ₂)	HPIC	solution
Carbon disulphide (CS ₂)	GC-MS, GC/FPD	gas
Hydrogene Sulphide (H ₂ S)	HPIC GC/FPD	solution solution
Ammonia (NH ₃)	Colourimetry HPIC Titration	solution solution solution
Antimony compounds	AAS or ICP	solution
Arsenic compounds	AAS or ICP	solution
Phosphorus	ICP	solution
Phosphates	Colourimetry HPIC	solution solution