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ICS 71.100.80

EAST AFRICAN STANDARD

Chemicals used for treatment of water intended for human consumption — Sodium disulfite

EAST AFRICAN COMMUNITY

Foreword

Development of the East African Standards has been necessitated by the need for harmonizing requirements governing quality of products and services in East Africa. It is envisaged that through harmonized standardization, trade barriers which are encountered when goods and services are exchanged within the Community will be removed.

In order to meet the above objectives, the EAC Partner States have enacted an East African Standardization, Quality Assurance, Metrology and Test Act, 2006 (EAC SQMT Act, 2006) to make provisions for ensuring standardization, quality assurance, metrology and testing of products produced or originating in a third country and traded in the Community in order to facilitate industrial development and trade as well as helping to protect the health and safety of society and the environment in the Community.

East African Standards are formulated in accordance with the procedures established by the East African Standards Committee. The East African Standards Committee is established under the provisions of Article 4 of the EAC SQMT Act, 2006. The Committee is composed of representatives of the National Standards Bodies in Partner States, together with the representatives from the private sectors and consumer organizations. Draft East African Standards are circulated to stakeholders through the National Standards Bodies in the Partner States. The comments received are discussed and incorporated before finalization of standards, in accordance with the procedures of the Community.

Article 15(1) of the EAC SQMT Act, 2006 provides that "Within six months of the declaration of an East African Standard, the Partner States shall adopt, without deviation from the approved text of the standard, the East African Standard as a national standard and withdraw any existing national standard with similar scope and purpose".

East African Standards are subject to review, to keep pace with technological advances. Users of the East African Standards are therefore expected to ensure that they always have the latest versions of the standards they are implementing.

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Introduction

In the preparation of this East African Standard, the following source was consulted extensively:

BS EN 12121:2005, *Chemicals used for treatment of water intended for human consumption — Sodium disulfite*

Assistance derived from this source and others inadvertently not mentioned is hereby acknowledged.

Draft for comments only — Not to be cited as East African Standard

Chemicals used for treatment of water intended for human consumption — Sodium disulfite

The European Standard EN 12121:2005 has the status of a
British Standard

ICS 71.100.80

National foreword

This British Standard is the official English language version of EN 12121:2005. It supersedes BS EN 12121:1999 which is withdrawn.

The UK participation in its preparation was entrusted to Technical Committee CII/59, Chemicals for drinking water treatment, which has the responsibility to:

- aid enquirers to understand the text;
- present to the responsible international/European committee any enquiries on the interpretation, or proposals for change, and keep UK interests informed;
- monitor related international and European developments and promulgate them in the UK.

A list of organizations represented on this committee can be obtained on request to its secretary.

The United Kingdom, as a member of CEN, is obliged to publish EN 12121 as a British Standard. However, attention is drawn to the fact that the United Kingdom voted against the approval of EN 12121. The main reasons for this disapproval were as follows.

- a) In the opinion of the UK committee, the limit on the sum of the contents of sodium sulfate and sodium chloride, quoted in 4.3, is unnecessary.
- b) No information has been provided on the preparation of test solutions or the calculation of results for the determinations of sulfate and chloride, in order to apply the methods referred to in 5.2.3 which are intended for the analysis of water.

Cross-references

The British Standards which implement international or European publications referred to in this document may be found in the *BSI Catalogue* under the section entitled "International Standards Correspondence Index", or by using the "Search" facility of the *BSI Electronic Catalogue* or of British Standards Online.

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Compliance with a British Standard does not of itself confer immunity from legal obligations.

Summary of pages

This document comprises a front cover, an inside front cover, the EN title page, pages 2 to 18, an inside back cover and a back cover.

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English version

Chemicals used for treatment of water intended for human consumption - Sodium disulfite

Produits chimiques utilisés pour le traitement de l'eau destinée à la consommation humaine - Disulfite de sodium

Produkt zur Aufbereitung von Wasser den menschlichen Gebrauch - Natriumdisulfid

This European Standard was approved by CEN on 12 May 2005.

CEN members are bound to comply with the CEN/CENELEC Internal Regulations which stipulate the conditions for giving this European Standard the status of a national standard without any alteration. Up-to-date lists and bibliographical references concerning such national standards may be obtained on application to the Central Secretariat or to any CEN member.

This European Standard exists in three official versions (English, French, German). A version in any other language made by translation under the responsibility of a CEN member into its own language and notified to the Central Secretariat has the same status as the official versions.

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Foreword

This European Standard (EN 12121:2005) has been prepared by Technical Committee CEN/TC 164 "Water supply", the secretariat of which is held by AFNOR.

This European Standard shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by December 2005, and conflicting national standards shall be withdrawn at the latest by December 2005.

This document supersedes EN 12121:1998.

Significant technical differences between this edition and EN 12121:1998 are as follows:

- deletion of this reference to EU Directive 80/778/EEC of July, 15 1980 in order to take into account the latest Directive in force (see [1]).

According to the CEN/CENELEC Internal Regulations, the national standards organizations of the following countries are bound to implement this European Standard: Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Slovakia, Slovenia, Spain, Sweden, Switzerland and United Kingdom.

EN 12121:2005 (E)

Introduction

In respect of potential adverse effects on the quality of water intended for human consumption, caused by the product covered by this European Standard :

- a) this European Standard provides no information as to whether the product may be used without restriction in any of the Member States of the EU or EFTA;
- b) it should be noted that, while awaiting the adoption of verifiable European criteria, existing national regulations concerning the use and/or the characteristics of this product remain in force.

NOTE Conformity with this European Standard does not confer or imply acceptance or approval of the product in any of the Member States of the EU or EFTA. The use of the product covered by this European Standard is subject to regulation or control by National Authorities.

1 Scope

This European Standard is applicable to sodium disulfite used for treatment of water intended for human consumption. It describes the characteristics of sodium disulfite and specifies the requirements and the corresponding test methods for sodium disulfite. It gives information on its use in water treatment. It also determines the rules relating to safe handling and use (see Annex B).

2 Normative references

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

EN ISO 3696, *Water for analytical laboratory use - Specification and test methods (ISO 3696:1987)*

ISO 3165, *Sampling of chemical products for industrial use - Safety in sampling*

ISO 3629, *Photography-Processing chemicals- Specifications for potassium metabisulfite*

ISO 5993, *Sodium hydroxide for industrial use - Determination of mercury content - Flameless atomic absorption spectrometric method*

ISO 6206, *Chemical products for industrial use - Sampling – Vocabulary*

ISO 6353-1, *Reagents for chemical analysis. Part 1: General test methods*

ISO 8213, *Chemical products for industrial use – Sampling techniques – Solid chemical products in the form of particles varying from powders to coarse lumps*

ISO 9297, *Water quality- Determination of chloride- Silver nitrate titration with chromate indicator (Mohr's method)*

ISO 22743 (in preparation) *Water quality – Determination of sulfates by continuous flow analysis (CFA)*

3 Description

3.1 Identification

3.1.1 Chemical name

Sodium disulfite.

3.1.2 Synonym or common names

Sodium metabisulfite, sodium pyrosulfite.

3.1.3 Relative molecular mass

190,10.

3.1.4 Empirical formula

$\text{Na}_2\text{S}_2\text{O}_5$.

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3.1.5 Chemical formula

Na₂S₂O₅.

3.1.6 CAS-Registry Number¹⁾

7681-57-4.

3.1.7 EINECS reference²⁾

231-673-0.

3.2 Commercial form

The product is a crystalline powder.

3.3 Physical properties

3.3.1 Appearance and odour

The product is a white, free-flowing fine crystalline powder with a slight odour of sulfur dioxide.

3.3.2 Density

The density of the product is 2,7 g/cm³ at 20 °C.

The bulk density is approximately 1,3 g/cm³ at 20 °C.

3.3.3 Solubility (in water)

The solubility of the product in water is 540 g/l at 20 °C.

3.3.4 Vapour pressure

Not applicable.

3.3.5 Boiling point at 100 kPa³⁾

Not applicable.

3.3.6 Crystallisation point

The product decomposes above 150 °C.

3.3.7 Specific heat

Not known.

1) Chemical Abstracts Service Registry Number.

2) European Inventory of Existing Commercial Chemical Substances.

3) 100 kPa = 1 bar.

3.3.8 Viscosity dynamic

Not applicable.

3.3.9 Critical temperature

Not applicable.

3.3.10 Critical pressure

Not applicable.

3.3.11 Physical hardness

Not applicable.

3.4 Chemical properties

The solution is weakly acid. The pH value of a saturated aqueous solution of sodium disulfite is between 4,5 and 5,0.

Sodium disulfite dissolves in water forming sodium hydrogen sulfite.

At elevated temperatures (> 150 °C) or on contact with acids sulfur dioxide is generated.

Sodium disulfite reacts violently with oxidizing agents ; e.g. with sodium hypochlorite or hydrogen peroxide.

4 Purity criteria**4.1 General**

This European Standard specifies the minimum purity requirements for sodium disulfite used for the treatment of water intended for human consumption. Limits are given for impurities commonly present in the product. Depending on the raw material and the manufacturing process other impurities may be present and, if so, this shall be notified to the user and when necessary to relevant authorities.

NOTE Users of this product should check the national regulations in order to clarify whether it is of appropriate purity for treatment of water intended for human consumption, taking into account raw water quality, required dosage, contents of other impurities and additives used in the products not stated in this product standard.

Limits have been given for impurities and chemical parameters where these are likely to be present in significant quantities from the current production process and raw materials. If the production process or raw materials leads to significant quantities of impurities, by-products or additives being present, this shall be notified to the user.

4.2 Composition of commercial product

The content of sodium disulfite shall not be less than a mass fraction of 95 % ($\text{Na}_2\text{S}_2\text{O}_5$).

The concentration of sodium disulfite shall be within ± 5 % of the manufacturer's declared value.

4.3 Impurities and main by-products

The sum of the content of sodium sulfate and sodium chloride shall not exceed a mass fraction of 5% of the commercial product.

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4.4 Chemical parameters

The product shall conform to the requirements specified in Table 1.

Table 1 — Chemical parameters

Parameter		Limit mg/kg of commercial product
Antimony (Sb)	max.	1
Arsenic (As)	max.	1
Cadmium (Cd)	max.	1
Chromium (Cr)	max.	1
Lead (Pb)	max.	5
Mercury (Hg)	max.	1
Nickel (Ni)	max.	1
Selenium (Se)	max.	1
NOTE Other chemical parameters and indicator parameters are not relevant in sodium disulfite because the raw materials used in the manufacturing process are free of them. For parametric values of sodium disulfite on trace metal content in drinking water, see [1].		

5 Test methods

5.1 Sampling

Observe the general recommendations of ISO 3165 and take account of ISO 6206.

Prepare the laboratory sample(s) required by the relevant procedure described in ISO 8213.

5.2 Analyses

5.2.1 General

All reagents shall be of a recognized analytical grade and the water used shall conform to grade 3 specified in EN ISO 3696.

5.2.2 Main product

5.2.2.1 General

The sodium disulfite content shall be determined by the method for potassium disulfite described in ISO 3629.

5.2.2.2 Principle

Sodium disulfite is oxidized with a fixed volume of iodine. The excess of added iodine is titrated with sodium thiosulfate. The determination includes other sulfites in addition to $\text{Na}_2\text{S}_2\text{O}_5$, therefore the second titration, as described in ISO 3629, is omitted for the purpose of this determination.

5.2.2.3 Calculation

The equation given in ISO 3629 for potassium disulfite has to be adapted for sodium disulfite.

The sodium disulfite content C_1 , expressed as a mass fraction in %, is given by the following equation:

$$C_1 = 4,76 \left(\frac{(100 \times c_3) - (V \times c_2)}{m} \right) \quad (1)$$

where:

- c_2 is the actual concentration, in moles per litre, of the sodium thiosulfate solution;
- c_3 is the actual concentration, in moles per litre, of the iodine solution;
- V is the volume, in millilitres, of the sodium thiosulfate solution used for the titration;
- m is the mass, in grams, of the test portion used for the titration;
- 4,76 is a conversion factor for the mass of sodium disulfite equivalent to 1 mole of iodine (i.e. 47,6) X the conversion factor for millilitres to litres (i.e. 0,001) X 100 (for percentage);
- 100 is the volume, in millilitres of iodine solution added for total oxidation.

5.2.3 Impurities**5.2.3.1 Sulfate**

The sulfate content shall be determined in accordance with ISO 22743

5.2.3.2 Chloride

The chloride content shall be determined in accordance with ISO 9297.

5.2.4 Chemical parameters

5.2.4.1 General

The content of chemical parameters shall be determined using the procedures specified in Table 2.

Table 2 — Procedures for the determination of chemical parameters

Element	Reference	Method	Wavelength nm	Flame
As	see 5.2.4.3	Hydride AAS	193,7	n.a.
Sb	see 5.2.4.3	Hydride AAS	217,6	n.a.
Cd	ISO 6353-1 GM 29 See 5.2.4.2	AAS	228,8	air-acetylene
Cr	ISO 6353-1 GM 29 See 5.2.4.2	AAS	357,8	air-acetylene
Pb	ISO 6353-1 GM 29 See 5.2.4.2	AAS	217,0 or 283,3	air-acetylene
Ni	ISO 6353-1 GM 29 See 5.2.4.2	AAS	232,0	oxidizing air-acetylene
Se	see 5.2.4.3	Hydride AAS	196,0	n.a.
Hg	in accordance with ISO 5993	flameless AAS	253,6	n.a.

AAS Atomic absorption spectrometry
n.a. not applicable.

5.2.4.2 Determination of cadmium (Cd), chromium (Cr), lead (Pb) and nickel (Ni)

5.2.4.2.1 General

The content of Cd, Cr, Pb and Ni shall be determined taking into account ISO 6353-1, modified as described in 5.2.4.2.2.

5.2.4.2.2 Procedure

The elements Cd, Cr, Pb and Ni shall be determined using the standard addition method. The reference solutions shall be made by spiking the sample with standard solutions, which contain step-wise increasing contents of the elements to be determined. The amount of internal standard to be added can be estimated from a preliminary investigation, determining roughly the element content of the test sample from simple calibration. The steps in which internal standards are added shall be at least as high as the estimated content of the test sample.

Carry out a blank determination by repeating the procedure using the same quantities of all reagents but omitting the test sample.

5.2.4.2.3 Expression of results

Determine the correlation line by plotting the measured absorbances of the spiked measurement solutions in relation to the element content. The spiked measurement solutions are produced by adding defined quantities of element to the measurement solution. They contain stepwise increasing contents of the elements to be determined.

Read the concentration of each element in the test solution by extrapolation of the correlation line to absorbance $A=0$ (see Figure 1). Similarly determine the element concentration of the blank solution (see Figure 2) and subtract from the result obtained for the test solution. Alternatively, the evaluation can be carried out by linear regression. Additional dilution steps shall be compensated in the calculation.

This interim result (y) expressed in micrograms per litre needs to be converted to give the final concentration according to the equation in 5.2.4.2.4.

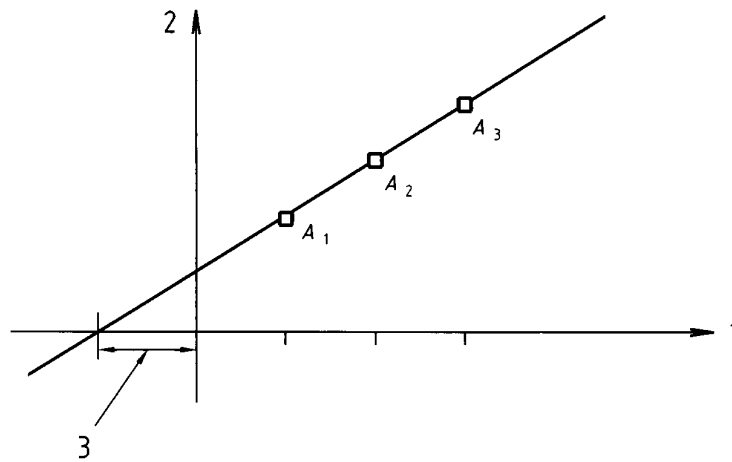
5.2.4.2.4 Calculation

From the interim result (y) determined (see 5.2.4.2.3) the content, C_3 , of each element in the laboratory sample, expressed in milligrams per kilogram of sodium disulfite is given by the following equation:

$$C_2 = \frac{y \times V \times 100 \times 1000}{m \times C_1} \tag{2}$$

where:

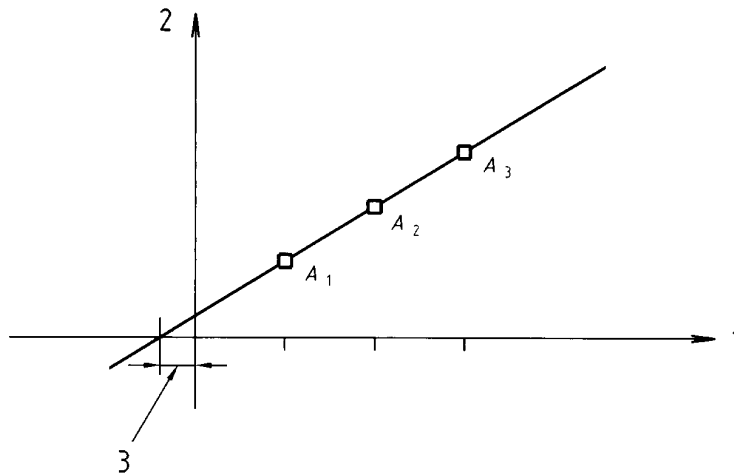
- y is the interim result (5.2.4.2.3);
- V is the volume, expressed in millilitres, of the test solution;
- m is the mass, expressed in grams, of the test portion;
- C_1 is the content, expressed in mass fraction in % of sodium disulfite (see 5.2.2.3).



Key

- 1 Concentration of added standard in micrograms per litre
- 2 Absorbance A
- 3 Concentration in the test solution in micrograms per litre
- A₁, A₂, A₃ spiking

Figure 1 — Calculation of the element concentration in the test solution



Key

- 1 Concentration of added standard in micrograms per litre
 - 2 Absorbance A
 - 3 Concentration in the blank solution in micrograms per litre
- A_1, A_2, A_3 spiking

Figure 2 — Calculation of the element concentration in the blank solution

5.2.4.3 Determination of arsenic (As), antimony (Sb) and selenium (Se)

5.2.4.3.1 Principle

The elements arsenic, antimony, and selenium are determined by hydride-atomic absorption spectrometry. The elements are reduced by sodium borohydride (NaBH_4) to form the hydrides. These volatile compounds flow through the heated measuring cuvette of an atomic absorption spectrometer where the content of the individual element is determined.

5.2.4.3.2 Reagents

5.2.4.3.2.1 Hydrochloric acid, high purity analytical grade, mass fraction 30 %, density $\rho = 1,15 \text{ g/ml}$.

5.2.4.3.2.2 Preliminary reduction agent

Dissolve 10 g sodium iodide and 100 g L-ascorbic acid, in 1 000 ml of water.

5.2.4.3.2.3 Reduction agent

Dissolve in water NaBH_4 and NaOH in concentrations specified in the manufacturer's handbook for the spectrometer.

5.2.4.3.2.4 Standard solution (100 $\mu\text{g/l}$ As, Sb or Se)

The standard solution shall be freshly prepared on the day of use by individual dilution of a stock solution. This stock solution with an As-, Sb- or Se- content of at least 1 mg/l shall be made by dilution of standard solutions of Se, As and Sb which are available from all major suppliers of laboratory chemicals. This stock solution shall be kept

in containers of tetrafluoroethylene-hexafluoropropylene copolymer (FEP), polytetrafluoroethylene (PTFE) or polyethylene (PE).

NOTE The stock solution should not be kept for longer than four weeks.

5.2.4.3.3 Apparatus

Ordinary laboratory apparatus and:

5.2.4.3.3.1 One one-mark volumetric flasks, 100 ml.

5.2.4.3.3.2 Nine one-mark volumetric flasks, 10 ml.

5.2.4.3.3.3 Pipettes 2,5 ml, 5ml, 10 ml and 20 ml.

5.2.4.3.3.4 Three one-mark volumetric flasks , 50 ml.

5.2.4.3.3.5 Micropipettes, volume adjustable to maximum 500 μ l.

5.2.4.3.3.6 Atomic absorption spectrometer with the measurement parameters specified in Table 2.

The width of the slit, the measuring time, flushing with argon before and after the measurement and the reaction time shall be adjusted in accordance with the manufacturer's instructions. The background compensation shall be activated for the measurement of As and Sb, but not for the measurement of Se.

5.2.4.3.4 Procedure

For As (procedure for Sb and Se in parentheses if different from As procedure):

Weigh a test portion of 1 g to the nearest 0,1 mg and transfer it into a 100 ml one-mark volumetric flask and make up to the mark at 20 °C with water. Pipette 10 ml (Sb, Se: 20 ml) of this solution into a 50 ml one-mark volumetric flask and add 5 ml of HCl and 5 ml of preliminary reduction agent. Do not add preliminary reduction agent to the flasks for Sb and Se determination. Allow 3 h for reaction to occur and fill to the mark with water. Pipette 2,5 ml (Sb, Se): 5 ml) of this solution into three 10 ml one-mark volumetric flasks labelled A, B, C. Add 0,8 ml of HCl (5.2.4.3.2.1) to each flask. For the purpose of internal calibration add those quantities of standard solutions as given in Table 3 to the flasks B and C. Using the spectrometer (5.2.4.3.3.6) carry out the measurement with the addition of the reduction agent (5.2.4.3.2.3) and the parameters of measurement in accordance with the manufacturer's instructions for the spectrometer.

Carry out a blank determination by repeating the procedure using the same quantities of all reagents but omitting the test sample.

Table 3 — Standard solution

	Volume of standard solution to be added		
	As	Sb	Se
Flask B	50 μ l	100 μ l	200 μ l
Flask C	100 μ l	200 μ l	500 μ l

5.2.4.3.5 Expression of results

See 5.2.4.2.3.

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5.2.4.3.6 Calculation

See 5.2.4.2.4.

6 Labelling - Transportation - Storage

6.1 Means of delivery

Sodium disulfite shall be delivered in plastics bags or plastic-lined paper bags.

In order that the purity of the product is not affected, the means of delivery shall not have been used previously for any different product or it shall have been specially cleaned and prepared before use.

6.2 Risk and safety labelling according to the EU Directives⁴⁾

The following labelling requirements shall apply to sodium disulfite at the date of the publication of this European Standard.

Symbols and indications of danger:

- Xi: irritant.

Nature of special risk attributed to dangerous substances:

- R 31: Contact with acids liberates toxic gas.
- R 36/37: Irritating to eyes and respiratory system.

Safety advice concerning dangerous substances:

- S2: Keep out of the reach of children.
- S26: In case of contact with eyes, rinse immediately with plenty of water and seek medical advice.

NOTE Annex I of the Directive 67/548/EEC on classification, packaging and labelling of dangerous substances and its amendments and adaptations in the European Union contains a list of substances classified by the EU. Substances not in this Annex I should be classified on the basis of their intrinsic properties according to the criteria in the Directive by the person responsible for the marketing of the substance.

6.3 Transportation regulations and labelling

Sodium disulfite is not listed under a UN number⁵⁾.

Sodium disulfite is not classified as a dangerous product for road, rail, sea and air transportation.

6.4 Marking

The marking shall include the following:

- name "sodium disulfite", trade name and grade;

⁴⁾ See [2].

⁵⁾ United Nations Number

- net mass;
- name and the address of supplier and/or manufacturer;
- statement "this product conforms to EN 12121".

6.5 Storage

6.5.1 Long term stability

The product is stable when stored in containers in a cool and dry place.

6.5.2 Storage incompatibilities

The product shall be kept away from acids, such as hydrochloric acid and sulfuric acid, to avoid the risk of sulfur dioxide evolution.

The product shall be kept away from oxidizing substances, such as sodium hypochlorite, hydrogen peroxide.

Annex A (informative)

General information on sodium disulfite

A.1 Origin

A.1.1 Raw materials

The sodium disulfite is manufactured from sulfur dioxide and sodium hydroxide.

A.1.2 Manufacturing process

Aqueous sodium hydroxide is treated with sulfur dioxide to produce sodium hydrogen bisulfite solution. This solution is cooled and sodium disulfite is removed by centrifuging. The commercial product is obtained after rapid drying.

A.2 Use

A.2.1 Function

Sodium disulfite is used as a reducing agent to remove excess chlorine, chlorine dioxide or ozone in the drinking water

A.2.2 Form in which it is used

As delivered and dissolved.

A.2.3 Treatment dose

The treatment dose depends on the content of oxidants in the water, e.g. for chlorine in water the stoichiometric dose is 1,34 mg of sodium disulfite for 1 mg of Cl_2 , but in practice an excess over the stoichiometric dose can be required.

A.2.4 Means of application

It is usually applied using a metering pump.

A.2.5 Secondary effects

The use of sodium disulfite increases the content of sodium and sulfate.

A.2.6 Removal of excess product

The excess product is removed by oxidation.

Annex B
(normative)

General rules relating to safety

B.1 Rules for safe handling and use

The supplier shall provide current safety instructions.

B.2 Emergency procedures

B.2.1 First aid

In case of contact with the skin wash immediately with plenty of water.

In case of contact with the eyes treat by irrigation with plenty of water, with the eyelids held open.

In case of ingestion seek immediately medical advice.

B.2.2 Spillage

Remove any spillage of solid product in avoiding the formation of dust.

Any remaining product can be flushed away with plenty of water to which an oxidizing agent is added.

B.2.3 Fire

The product is not combustible.

Decomposition can be caused by fire with release of sulfur dioxide. Wear suitable respiratory equipment.

There are no restrictions on extinguishing media in fire situations.

Bibliography

- [1] 98/83/EC: Council Directive of 3 November 1998 on the quality of water intended for human consumption
- [2] 67/548/EEC: Council Directive of 27th June 1967 on the approximation of the laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances and its amendments and adaptations

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