
**Copper, lead, zinc and nickel
concentrates — Sampling of slurries**

*Concentrés de cuivre, de plomb, de zinc et de nickel — Échantillonnage
des schlamms*



PDF disclaimer

This PDF file may contain embedded typefaces. In accordance with Adobe's licensing policy, this file may be printed or viewed but shall not be edited unless the typefaces which are embedded are licensed to and installed on the computer performing the editing. In downloading this file, parties accept therein the responsibility of not infringing Adobe's licensing policy. The ISO Central Secretariat accepts no liability in this area.

Adobe is a trademark of Adobe Systems Incorporated.

Details of the software products used to create this PDF file can be found in the General Info relative to the file; the PDF-creation parameters were optimized for printing. Every care has been taken to ensure that the file is suitable for use by ISO member bodies. In the unlikely event that a problem relating to it is found, please inform the Central Secretariat at the address given below.

This document is a preview generated by EVS



COPYRIGHT PROTECTED DOCUMENT

© ISO 2010

All rights reserved. Unless otherwise specified, no part of this publication may be reproduced or utilized in any form or by any means, electronic or mechanical, including photocopying and microfilm, without permission in writing from either ISO at the address below or ISO's member body in the country of the requester.

ISO copyright office
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

Published in Switzerland

Contents

Page

Foreword	v
1 Scope	1
2 Normative references	1
3 Terms and definitions	2
4 Principles of sampling slurries	2
4.1 General	2
4.2 Sampling errors	3
4.2.1 General	3
4.2.2 Preparation error, PE	4
4.2.3 Delimitation and extraction errors, DE and EE	4
4.2.4 Weighting error, WE	6
4.2.5 Periodic quality-fluctuation error, QE ₃	6
4.3 Sampling and total variance	6
4.3.1 Sampling variance	6
4.3.2 Total variance	6
4.3.3 Sampling-stage method of estimating sampling and total variance	8
4.3.4 Simplified method of estimating sampling and total variance	9
4.3.5 Interleaved sample method of measuring total variance	10
5 Establishing a sampling scheme	11
6 Minimization of bias and unbiased increment mass	16
6.1 Minimization of bias	16
6.2 Volume of increment for falling-stream samplers to avoid bias	17
7 Number of increments	17
7.1 General	17
7.2 Simplified method	18
8 Minimum mass of solids contained in lot and sub-lot samples	18
8.1 Minimum mass of solids in lot samples	18
8.2 Minimum mass of solids in sub-lot samples	18
8.3 Minimum mass of solids in lot and sub-lot samples after size reduction	18
9 Time-basis sampling	19
9.1 General	19
9.2 Sampling interval	19
9.3 Cutters	19
9.4 Taking of increments	19
9.5 Constitution of lot or sub-lot samples	20
9.6 Division of increments and sub-lot samples	20
9.7 Division of lot samples	20
9.8 Number of cuts for division	20
10 Stratified random sampling within fixed time intervals	20
11 Mechanical sampling from moving streams	21
11.1 General	21
11.2 Design of the sampling system	21
11.2.1 Safety of operators	21
11.2.2 Location of sample cutters	21
11.2.3 Provision for duplicate sampling	21
11.2.4 System for checking the precision and bias	21
11.2.5 Avoiding bias	22

11.3	Slurry sample cutters	22
11.3.1	General.....	22
11.3.2	Falling-stream cutters	23
11.3.3	Cutter velocities	23
11.4	Mass of solids in increments.....	23
11.5	Number of primary increments	23
11.6	Routine checking	23
12	Manual sampling from moving streams	24
12.1	General.....	24
12.2	Choosing the sampling location	24
12.3	Sampling implements.....	25
12.4	Mass of solids in increments.....	25
12.5	Number of primary increments	25
12.6	Sampling procedures	25
13	Sampling of stationary slurries	26
14	Sample preparation	26
14.1	General.....	26
14.2	Sample division.....	26
14.3	Sample grinding.....	26
14.4	Chemical analysis samples	26
14.5	Physical test samples	27
15	Packing and marking of samples.....	27
Annex A	(normative) Sampling-stage method for estimating sampling and total variance	28
Annex B	(informative) Examples of correct slurry sampling devices	34
Annex C	(informative) Examples of incorrect slurry sampling devices	37
Annex D	(normative) Manual sampling implements.....	41
Bibliography	42

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 11794 was prepared by Technical Committee ISO/TC 183, *Copper, lead, zinc and nickel ores and concentrates*.

This document is a preview generated by EVS

Copper, lead, zinc and nickel concentrates — Sampling of slurries

WARNING — This International Standard may involve hazardous materials, operations and equipment. It is the responsibility of the user of this International Standard to establish appropriate health and safety practices and determine the applicability of regulatory limitations prior to use.

1 Scope

This International Standard sets out the basic methods for sampling particulate material that is mixed with a liquid, usually water, to form a slurry. In industry and in the mining and mineral processing literature, slurry is also referred to as pulp, but this term is not used in this International Standard. At very high ratios of fine particulate solids to liquids where material assumes a soft plastic form, the mixture is correctly termed as a paste. Sampling of pastes is not covered in this International Standard.

The procedures described in this International Standard apply to sampling of particulate materials that are transported in moving streams as slurries, but not pressurized slurries. These streams may fall freely or be confined in pipes, launders, flumes, sluices, spirals or similar channels. Sampling of slurries in stationary situations, such as a settled or even a well-stirred slurry in a holding vessel or dam, is not recommended and is not covered in this International Standard.

This International Standard describes procedures that are designed to provide samples representative of the slurry solids and particle-size distribution of the slurry under examination. After draining the slurry sample of fluid and measuring the fluid volume, damp samples of the contained particulate material in the slurry are available for drying (if required) and measurement of one or more characteristics in an unbiased manner and with a known degree of precision. The characteristics are measured by chemical analysis, physical testing or both.

The sampling methods described are applicable to slurries that require inspection to verify compliance with product specifications, determination of the value of a characteristic as a basis for settlement between trading partners or estimation of a set of average characteristics and variances that describes a system or procedure.

Provided that flow rates are not too high, the reference method against which other sampling procedures are compared is one where the entire stream is diverted into a vessel for a specified time or volume interval. This method corresponds to the stopped-belt method described in ISO 12743.

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.

ISO 12743, *Copper, lead, zinc and nickel concentrates — Sampling procedures for determination of metal and moisture content*

ISO 12744, *Copper, lead, zinc and nickel concentrates — Experimental methods for checking the precision of sampling*

ISO 13292, *Copper, lead, zinc and nickel concentrates — Experimental methods for checking the bias of sampling*

ISO 20212, *Copper, lead, zinc and nickel sulfides — Sampling procedures for ores and smelter residues*

3 Terms and definitions

For the purposes of this document, the terms and definitions given in ISO 12743, ISO 12744, ISO 13292 and ISO 20212 apply.

4 Principles of sampling slurries

4.1 General

In this International Standard, a slurry is defined as “any fluid mixture of a solid of nominal top size < 1 mm that is mixed with water, which is frequently used as a convenient form to handle solids in bulk”. Slurry flows are found in many mineral processing plants, with the water and entrained solids mixture being transported through the plant circuits by means of pumps and pipelines and under gravity in sluices, flumes and launders. In a number of operations, ore is transported to the mill in slurry form, and in others concentrates are transported long distances in slurry pipelines. Tailings from wet plants are also discharged as slurries through pipelines to the tailings dam. In many of these operations, collection of increments at selected sample points is required for evaluation of the particulate material in the slurry.

A lot sample is constituted from a set of unbiased primary increments from a lot. The sample container is weighed immediately after collection and combination of increments to avoid water loss by evaporation or spillage. Weighing is necessary to determine the percentage of solids by mass in the slurry sample. The sample may then be filtered, dried and weighed. Alternatively, the sample may be sealed in plastic bags after filtering for transport and drying at a later stage. The liquid removed during filtration should be retained if it needs to be analysed.

Test samples are prepared from samples after filtering and drying. Test portions may then be taken from the test sample and analysed using an appropriate and properly calibrated analytical method or test procedure under prescribed conditions.

The objective of the measurement chain is to determine the characteristic of interest in an unbiased manner with an acceptable and affordable degree of precision. The general sampling theory, which is based on the additive property of variances, can be used to determine how the variances of sampling, sample preparation and chemical analysis or physical testing propagate and hence determine the total variance for the measurement chain. This sampling theory can also be used to optimize manual sampling methods and mechanical sampling systems.

If a sampling scheme is to provide representative samples, all parts of the slurry in the lot must have an equal opportunity of being selected and appearing in the sample for testing. Hence, slurries are to be sampled in such a manner that all possible primary increments in the set into which the slurry can be divided have the same probability of being selected. Any deviation from this basic requirement can result in bias. A sampling scheme having incorrect selection techniques, i.e. with non-uniform selection probabilities, cannot be relied upon to provide representative samples.

Sampling of slurries should preferably be carried out by systematic sampling on a time basis (see Clause 9). If the slurry flow rate and the solids concentration vary with time, the slurry volume and the mass of dry solids for each increment will vary accordingly. It needs to be shown that no systematic error (bias) is introduced by periodic variation in quality or quantity, where the proposed sampling interval is approximately equal to a multiple of the period of variation in quantity or quality. Otherwise, stratified random sampling should be used (see Clause 10).