
**Determination of particle size distribution
by centrifugal liquid sedimentation
methods —**

Part 2:
Photocentrifuge method

*Détermination de la distribution granulométrique par les méthodes de
sédimentation centrifuge dans un liquide —*

Partie 2: Méthode photocentrifuge



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.

STANDARDSISO.COM : Click to view the full PDF of ISO 13318-2:2001

© ISO 2001

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.ch
Web www.iso.ch

Printed in Switzerland

Contents

Page

Foreword.....	iv
Introduction.....	v
1 Scope	1
2 Normative references	1
3 Terms, definitions and symbols.....	2
3.1 Terms and definitions	2
3.2 Symbols	2
4 Principle.....	2
5 Apparatus	3
6 Sampling.....	5
7 Preparation.....	5
7.1 Sample preparation	5
7.2 Temperature	5
7.3 Dispersion	5
8 Procedure	5
8.1 Line-start methods	5
8.2 Homogeneous technique.....	6
9 Tests in duplicate and validation	7
9.1 Tests in duplicate	7
9.2 Validation.....	7
10 Calculation of results	7
10.1 General.....	7
10.2 Calculation of particle size	7
10.3 Calculation of cumulative mass percentage	7
11 Reporting of results.....	8
Annex A (informative) Worked example	10
Annex B (informative) Extinction curve, example for titanium dioxide.....	14
Annex C (informative) Effect of radial dilution.....	15

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 3.

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 part of ISO 13318 may be the subject of patent rights. ISO shall not be held responsible for identifying any or all such patent rights.

ISO 13318-2 was prepared by Technical Committee ISO/TC 24, *Sieves, sieving and other sizing methods*, Subcommittee SC 4, *Sizing by methods other than sieving*.

ISO 13318 consists of the following parts, under the general title *Determination of particle size distribution by centrifugal liquid sedimentation methods*:

- *Part 1: General principles and guidelines*
- *Part 2: Photocentrifuge method*
- *Part 3: Centrifugal X-ray method*

Annexes A to C of this part of ISO 13318 are for information only.

Introduction

The sample suspension in a photocentrifuge may be contained in a cuvet or a disc. Sample concentration is determined by changes in a light signal monitored at a known radius. The cuvet photocentrifuge can only be run in the homogeneous mode whereas the disc photocentrifuge may be run in either the homogeneous or the line-start mode. Some systems permit the coarse end of the distribution to be measured in a gravitational mode and the fine end in the centrifugal mode. The use of light to determine particle size distribution requires a calibration factor to be applied as the particle size approaches the wavelength of the light, due to the inapplicability of the laws of geometric optics.

STANDARDSISO.COM : Click to view the full PDF of ISO 13318-2:2001

Determination of particle size distribution by centrifugal liquid sedimentation methods —

Part 2: Photocentrifuge method

WARNING — This part of ISO 13318 may involve hazardous materials, operations and equipment. This part of ISO 13318 does not purport to address all of the safety problems associated with its use. It is the responsibility of the user of this part of ISO 13318 to establish appropriate safety and health practices and determine the applicability of the regulatory limitations prior to its use.

1 Scope

This part of ISO 13318 covers methods for determining the particle size distribution of particulate materials by means of centrifugal sedimentation in a liquid. Solids concentrations are determined by the transmission of a light beam. The resulting signal enables conversion to a particle size distribution.

The method of determining the particle size distribution described in this part of ISO 13318 is applicable to powders that can be dispersed in liquids, powders that are present in slurry form and some emulsions. Typical particle size range for analysis is from about 0,1 μm to 5 μm . The method is applicable to powders in which all particles have the same density and comparable shapes and do not undergo chemical or physical change in the suspension liquid. It is usually necessary that the particles have a density higher than that of the liquid.

2 Normative references

The following normative documents contain provisions which, through reference in this text, constitute provisions of this part of ISO 13318. For dated references, subsequent amendments to, or revisions of, any of these publications do not apply. However, parties to agreements based on this part of ISO 13318 are encouraged to investigate the possibility of applying the most recent editions of the normative documents indicated below. For undated references, the latest edition of the normative document referred to applies. Members of ISO and IEC maintain registers of currently valid International Standards.

ISO 13318-1, *Determination of particle size distribution by centrifugal liquid sedimentation methods — Part 1: General principles and guidelines*

ISO 14887, *Sample preparation — Dispersing procedures for powders in liquids*

3 Terms, definitions and symbols

3.1 Terms and definitions

For the purposes of this part of ISO 13318 the terms and definitions of ISO 13318-1 apply.

3.2 Symbols

For the purposes of this this part of ISO 13318 the symbols of ISO 13318-1 and the following apply:

- D optical density
- E_i extinction coefficient for a particle of diameter x_i
- G constant dependent upon the geometry of the system, the dimensions of the light beam and on the shape of the particles
- I transmission of the emergent light beam, at the time t , after the start of sedimentation
- I_0 transmission of the emergent light beam when no particles are present
- M distance from rotation axis to measurement zone (mm)
- n_i number of particles of diameter x_i in the beam
- R distance from rotation axis to centrifuge wall, inner disc radius (mm)
- S distance from rotation axis to liquid-air interface of sample (mm)
- x_0 diameter of the smallest particle in the light beam (μm)
- x_{St} diameter of the largest particle in the light beam i.e. the Stokes diameter (μm)

4 Principle

A stable, finely collimated beam of light passes through a spinning disc or cuvet and sedimenting sample and is detected at a known radius. Light rays, typically from either a white light source (e.g. incandescent bulb) or a monochromatic coherent source (e.g. laser), pass through the suspension and are detected by a photodiode or photomultiplier. The disc photocentrifuge can be operated in the line-start or homogeneous mode whereas the cuvet photocentrifuge can be operated only in the homogeneous mode. The signal of the light beam is monitored over the analysis time. The mass percentage of sample present in the beam is determined by calculating the ratio of the light transmission signal, by use of a clear dispersing liquid, to the light transmission signal with the sample present.

In the line-start mode the disc initially contains clear fill liquid to give maximum light transmission. Then the sample is injected as a thin layer on top of the spinning fill liquid and begins to settle outward radially. When the largest particles present reach the light beam the light transmission decreases, returning to the original transmission value when the smallest particle present passes through the beam. A buffer layer is usually injected over the fill liquid to prevent suspension breaking through the interface in a phenomenon known as "streaming".

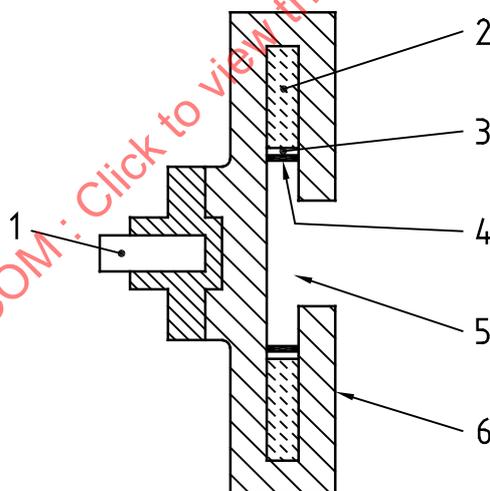
5 Apparatus

5.1 Disc photocentrifuge, with a chamber consisting of a hollow disc with an entry port coaxial with the axis of rotation (see Figure 1). Typically this is mounted vertically, or at a small angle to the vertical, on to the shaft of an electric motor with a digitally variable speed typically between $500 \text{ r}\cdot\text{min}^{-1}$ and $15\,000 \text{ r}\cdot\text{min}^{-1}$. A white light source and detector assembly measures transmittance through the suspension as a function of time. The instrument can be used in either a line-start or homogeneous mode. Extinction coefficient corrections need to be applied for the breakdown in the laws of geometric optics for both line-start and homogeneous modes. Additionally a correction is required for radial dilution effects when the homogeneous mode is used. Software is provided with commercial equipment to convert the data directly into size distributions in the form of tables or graphs of cumulative mass percentage versus particle size.

5.2 Cuvet photocentrifuge, in which the disc is replaced with a rectangular cell containing a homogeneous suspension (see Figure 2). Corrections need to be made for both radial dilution and light scattering effects as described in ISO 13318-1. Cuvet photocentrifuges can typically be run in both the gravitational and centrifugal modes. Additionally, some systems may offer a gradient mode permitting the centrifuge to accelerate throughout the analysis in order to reduce the measurement time.

5.3 Ancillary apparatus, consisting of:

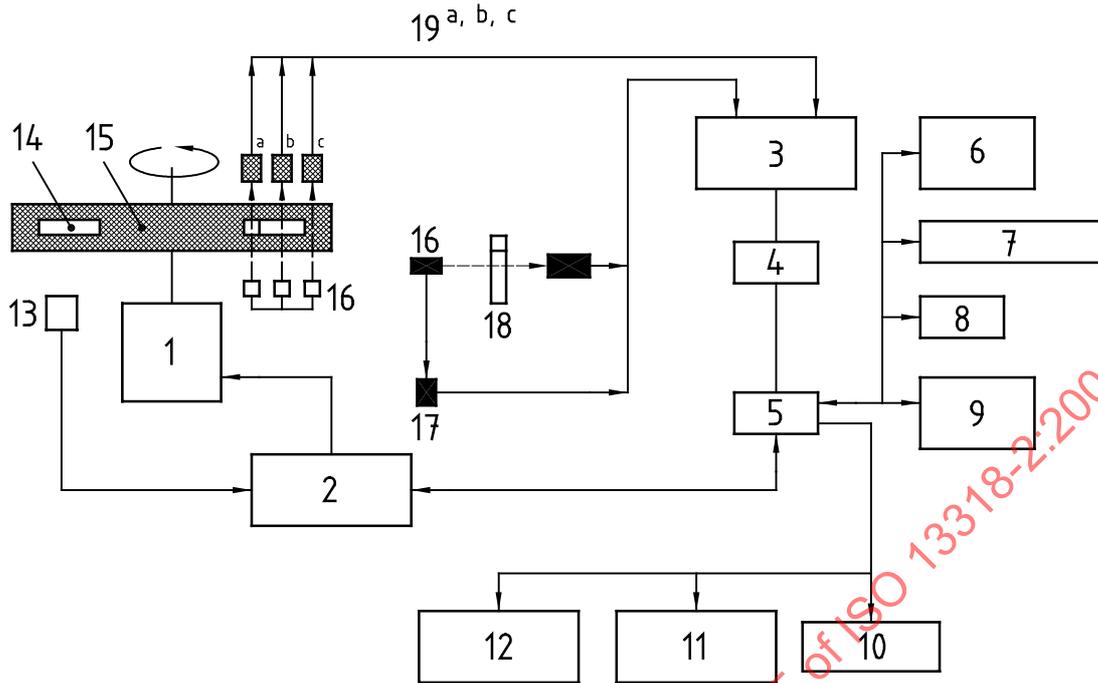
- dispersing vessel e.g. glass beaker or bottle, of appropriate dimensions;
- flexible spatula;
- ultrasonic bath or probe, a bottle shaker or high speed mechanical stirrer capable of rotating at $500 \text{ r}\cdot\text{min}^{-1}$ to $1\,000 \text{ r}\cdot\text{min}^{-1}$.



Key

- 1 Motor shaft
- 2 Spin fluid
- 3 Buffer layer
- 4 Suspension
- 5 Entry port
- 6 Polymethylmethacrylate disc

Figure 1 — Side view of the disc of a disc photocentrifuge



Key

- 1 Motor
- 2 Motor control
- 3 Signal process
- 4 ADC
- 5 CPU
- 6 Date time
- 7 Analysis parameters
- 8 RPM
- 9 Key board
- 10 Printer
- 11 Analog interface
- 12 Computer interface
- 13 Photo sensor RPM
- 14 Centrifugal cell
- 15 Rotating disc
- 16 LED
- 17 Photocell (reference)
- 18 Photocell (sample)
- 19 Photocells

- a Synchro-signal (reference)
- b Analogue signal
- c Synchro-signal (sample)

Figure 2 — Schematic diagram of a typical cuvet photocentrifuge

6 Sampling

For sampling see ISO 13318-1.

7 Preparation

7.1 Sample preparation

An analysis sample shall be prepared as described in ISO 13318-1. The volume and concentration required depends upon the volume of the centrifuge disc (or cuvet), the sensitivity of the optical-electronic system and whether the line-start (disc only) or homogeneous method is to be used. In general, lower concentrations are required than for other sedimentation methods. A concentration typically less than 0,25 % *m/v* and providing an attenuation preferably in the 20 % to 30 % range compared to the spin fluid without sample is required.

7.2 Temperature

The temperature of the spin fluid (line-start method) or suspension (homogeneous method) shall be determined and recorded before and after analysis in accordance with ISO 13318-1. The liquid viscosity and liquid density shall be recorded for the spin fluid or suspension at the temperature of the analysis. The temperature shall be maintained in accordance with ISO 13318-1.

7.3 Dispersion

For dispersion see ISO 13318-1 and ISO 14887.

8 Procedure

8.1 Line-start methods

8.1.1 General

The line-start method is one in which a layer of sample suspension is deposited on a spinning fill liquid i.e. all particles are considered to have started to sediment from the same initial radius. However the sample suspension has a tendency to penetrate the surface of the spin liquid as globules of suspension which then "stream" rapidly through the spin liquid. The streaming effect is due to an uneven outward diffusion of the sample suspension. On injection of the sample suspension the particles can become concentrated by surface tension effects at the interface. This leads to a region of high density above one of lower density and causes a bulk transfer of concentrated suspension into the spin fluid. Streaming is particularly prone to happen if the sample suspension is injected directly on to the spin fluid (two-layer method); consequently the two-layer method is in general replaced by the three-layer method (see 8.1.2) due to its applicability to a wider range of sample types. The following methods all attempt to cushion the hydrodynamic shock experienced by the particles as they enter the spin fluid and thus obtain a smooth transition and laminar flow.

Hydrodynamically stable sedimentation can generally be achieved by using the three-layer method, the buffered line or the external gradient method.

8.1.2 Three-layer method

Partially fill the centrifuge disc with a known volume of clear liquid (the spin fluid, typically approximately 15 ml) and allow the disc to attain the required running speed. Set up the light source at the required sampling radius (M). A steady baseline represents maximum transmission (I_0). Introduce into the disc a small volume of buffer liquid, typically 0,5 ml or 1 ml, that is of lower density than the spin fluid and determine the vortex radius (S). This can be done from a knowledge of the dimensions of the disc and the total volume of liquid introduced. Introduce the required volume of suspension (typically 0,25 ml of a 0,25 % *m/v* concentration) into the disc and activate the timer.

The expectation is that any rapid turbulence caused by the injection of the sample suspension is restrained to the intermediate (buffer) layer and be used to effect good sample mixing. The spin fluid might typically consist of 15 ml of 10 % (v/v) aqueous glycerol on which is floated 0,5 ml of water as the buffer layer followed by 0,25 ml of sample suspension. If streaming persists it may frequently be eliminated by using much smaller volumes of buffer liquid and sample suspension e.g. 0,1 ml.

In order for a line-start to be valid, the band of suspension has to be thin relative to the distance from the injection radius (S) to the radius of the measurement zone (M). It is recommended that the thickness of the suspension layer be greater than 5 % of ($M-S$).

As the particles are centrifuged outwards they attenuate the light beam and the changes in transmission are recorded. Run the experiment until the spin fluid is once again clear and the trace has returned to the baseline. For samples containing a large number of fine particles it may take a considerable time for the trace to return to the baseline. In these circumstances it may be expedient to terminate the run at an earlier time, e.g., when the signal has fallen to 10 % of the maximum deflection. A second run may then be made at a higher centrifuge speed to cover the fine end of the size range.

8.1.3 Buffered line-start method

This is a variation of the three-layer method. Interpose a thin layer of the suspension medium liquid, rather than a different buffer layer medium, between the spin liquid and the suspension layer. After the addition of the buffer layer apply transient deceleration to the disc so that the buffer is mixed with the outermost layer of the spin liquid. Inject the suspension sample on to the surface of the mixed layer once the disc has regained synchronous speed. Proceed as in 8.1.2.

8.1.4 External gradient method

0,2 ml of methanol is injected into a dry, non-rotating disc. Without delay the disc is brought to the required speed and 15 ml of distilled or de-ionized water is injected into the spinning disc. Immediately 0,1 ml, or less, of dodecane is injected into the disc to minimise evaporative cooling. Wait five minutes to enable the liquids and disc to approach thermal equilibrium and to permit the less dense methanol to rise through the water. Although methanol and water are miscible in all proportions, under the constant centrifugal force the less dense methanol will rise towards the meniscus to produce the required gradient. 0,2 ml of sample, of concentration about 0,1 % by volume, is injected and the analysis is started. The analysis temperature is recorded at the end of the run.

8.2 Homogeneous technique

Fill the centrifuge disc with pure suspending liquid, spin to the required speed and record the I_0 value. Empty the disc. Spin the disc to the required speed and pour in (or inject using a hypodermic syringe) the sample suspension through the entry port as quickly as possible. When the sample addition is complete activate the timer without delay.

The principle for the cuvet photocentrifuge is outlined in Figure 2. Light from a high intensity source is typically focused in two directions, one through a gravitational sample port and one through the measurement plane of the centrifuge. The collimated beams are filtered with narrow band pass filters. In the centrifugal mode two cells are used. The signal difference between the reference cell, containing clean suspension liquid, and the suspension cell are recorded once for each revolution. The reference cell is filled with clean liquid and inserted followed by the suspension cell filled with well-dispersed suspension. The centrifuge is switched on and typically this will initiate the timer to record the data.

9 Tests in duplicate and validation

9.1 Tests in duplicate

For tests in duplicate see ISO 13318-1.

9.2 Validation

For validation see ISO 13318-1. However, use reference materials with caution. Note that where it is intended to apply an extinction coefficient correction that this is a function of the sample material as well as the particle size and that the BCR and NIST reference materials are not all of the same chemical composition.

10 Calculation of results

10.1 General

The photocentrifuge measures the optical density of a suspension as a function of time from the commencement of the analysis. For the homogeneous mode, the mass percent undersize is obtained (if geometric optics apply) by normalizing the optical density, after correcting for radial dilution effects (see ISO 13318-1).

10.2 Calculation of particle size

Stokes diameters shall be calculated in accordance with ISO 13318-1.

10.3 Calculation of cumulative mass percentage

10.3.1 General

The light obscured by particles suspended in the path of a parallel beam of light is related to the concentration by the equation:

$$\ln\left(\frac{I_0}{I}\right) = G \sum_{i=0}^{St} n_i x_i^2 E_i \quad (1)$$

E_i is defined as the ratio of light obscured by a particle of diameter (x_i) to the light which would be obscured if the laws of geometric optics were valid for the system under consideration. If, for a given system using monochromatic light, the ratio of particle diameter to the incident wave length λ , is greater than about 100, the value of E_i is essentially constant. However for smaller ratios, E_i becomes a complex function of x_i/λ , (the relative refractive index of the particle with respect to that of the suspension medium), the shape of the particle and the geometry of the measuring device. Although fluctuations in E_i can be minimized by using a white light source in conjunction with a wide-angle acceptance detection receiver, the method is not absolute for particles whose diameter approximates to the wavelength of the incident light. Unless, therefore, the values of E_i corresponding to different diameters have either been calculated or measured experimentally for the system under test, results obtained by the method can only be regarded as having an unknown accuracy. The relationship between the time of sedimentation and the attenuation of the light beam is a characteristic of the sedimenting suspension and can be used for comparative investigations or quality control purposes.

10.3.2 Line-start technique (mass %)

For the line-start technique it is assumed that the particles present in the measurement zone at time t are of diameter x_{St} since the particles start from the same radius. The beam of light has a finite width and particles in the beam will constitute a size range proportional to x_{St} . The optical density, $D_t = -\log_{10}(I_t/I_0)$, at time t is therefore proportional to $n_t x_{St}^3$ (for geometric optics, where E_i does not vary with x) since the range of summation of equation (1) is proportional to $x_{St} t$.

A table is constructed showing variations of optical density with x_{St} at convenient time intervals. It is not necessary for the transmission data to be in absolute units and logarithms to base 10 can be used since the results are eventually to be expressed as mass percentages. Values of $\log_{10}(I_0/I)$, are then plotted against x_{St} . If it is assumed that E_i and the particle shape coefficient G are constant then this graph represents the mass frequency curve which can be integrated by areas to give the cumulative percentage by mass undersize data (see ISO 13318-1).

10.3.3 Homogeneous technique (mass %)

The photocentrifuge measures the optical density of a suspension against time from commencement of analysis. For the homogeneous mode, normalizing the ordinate (optical density), after correcting for the radial dilution effect, and converting the abscissa to Stokes diameter generates a curve of optical density against Stokes diameter. Data in this form may be suitable for comparison purposes; otherwise corrections (e.g. Mie) may be used to correct for the breakdown in the laws of geometric optics with particle size. The measured surface concentration (nx^2) is determined using equation (1). Correction is made for radial dilution (see ISO 13318-1) to give the frequency undersize by surface. The frequency undersize by mass is obtained by summing the product of $F(\text{surface})$ and x_{St} and equating the total to 100 % (see Table A.1).

11 Reporting of results

The report shall conform with the requirements set out in ISO 13318-1. Results shall be presented as Stokes diameter versus cumulative distribution by mass reported to the nearest 0.1 %. In the case of a plot, the diameters shall be placed on the abscissa and the cumulative mass percentage on the ordinate.

The test report shall contain the following information:

- a) reference to this part of ISO 13318, i.e ISO 13318-2;
- b) name of the testing establishment;
- c) date of the test;
- d) unique report identification;
- e) operator's identity;
- f) instrument type used;
- g) mode of operation (e.g. line-start, homogeneous);
- h) sample identification;
- i) sample material;
- j) sample density (and mass, if applicable);
- k) suspending liquid, its temperature, its density, its viscosity and, where applicable, its volume;
- l) Mie correction applied or not applied;
- m) sample volume and concentration injected;
- n) dispersing agent and concentration of agent used;
- o) method of dispersion of the suspension, including dispersion time;
- p) buffer layer, and where applicable, the type and volume;

- q) centrifuge speed;
- r) light source;
- s) any other operation not specified in this part of ISO 13318 and which could have an influence on the results.

The following instrument characteristics, typically determined by the manufacturer and preprogrammed into the equipment software, may be optionally reported if available:

- measurement radius (M);
- inner disc radius (R);
- inner disc thickness (used to calculate vortex radius, S).

STANDARDSISO.COM : Click to view the full PDF of ISO 13318-2:2001

Annex A (informative)

Worked example

Worked example for a photocentrifuge analysis in line-start mode:

Reference to the International Standard	ISO 13318-2
Name of the testing establishment	MAL
Date of the test	2000-09-23
Unique report identification	Carbon 123
Operator's identity	JM
Instrument type used	Model P123
Operation mode	Line-start
Sample identification	C-B No.1
Sample material	Carbon black
Sample density	1 860 kg·m ⁻³
Spin liquid	Water
Temperature (spin fluid)	293,15 K
Liquid density	1 065 kg·m ⁻³
Liquid viscosity	2,10 mPa·s
Spin liquid volume	10 ml
Mie correction	Not applied in Table A.1. Applied in Figure A.2
Sample volume and concentration	0,2 ml, 0,25 % <i>m/v</i>
Dispersing agent	0,05 % <i>m/v</i> sodium polymetaphosphate in distilled water
Dispersion method	1 min ultrasonic probe (75 W) in water-cooled jacketed container
Buffer layer	20/80 glycerol/water ¹⁾
Buffer layer volume	1 ml
Centrifuge speed	8 000 r·min ⁻¹

1) Distilled water containing 0,05 % *m/v* sodium polymetaphosphate.

Measurement radius (M)	4,815 cm
Inner disc radius (R)	5,073 cm
Inner disc thickness	0,623 4 cm
Light source	LED 560 nm

Remarks:

$$\text{Vortex radius, } S = \sqrt{\frac{\pi(5,073)^2(0,623\ 4) - 10}{\pi 0,623\ 4}} = 4,542\ \text{cm}$$

Column 1 of Table A.1 gives the elapsed time from the commencement of the run and column 2 gives the measured optical density. These data are presented graphically in Figure A.1.

STANDARDSISO.COM : Click to view the full PDF of ISO 13318-2:2001