
Cleanrooms and associated controlled environments —

**Part 9:
Assessment of surface cleanliness for
particle concentration**

Salles propres et environnements maîtrisés apparentés —

*Partie 9: Évaluation de la propreté des surfaces en fonction de la
concentration de particules*

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Contents

	Page
Foreword.....	iv
Introduction.....	v
1 Scope.....	1
2 Normative references.....	1
3 Terms and definitions.....	1
4 Abbreviated terms.....	2
5 The surface cleanliness level assessment system.....	3
5.1 ISO-SCP grading level format.....	3
5.2 Designation.....	6
5.3 General information on surface cleanliness levels of particle concentration.....	6
6 Demonstration of conformity.....	6
6.1 Principle.....	6
6.2 Testing.....	6
6.3 Test report.....	7
Annex A (informative) Surface characteristics.....	9
Annex B (informative) Descriptor for specific particle size ranges.....	12
Annex C (informative) Parameters influencing the SCP grading level assessments.....	15
Annex D (informative) Measurement methods for determining surface cleanliness by particle concentration.....	17
Bibliography.....	26

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.

The procedures used to develop this document and those intended for its further maintenance are described in the ISO/IEC Directives, Part 1. In particular, the different approval criteria needed for the different types of ISO documents should be noted. This document was drafted in accordance with the editorial rules of the ISO/IEC Directives, Part 2 (see www.iso.org/directives).

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. Details of any patent rights identified during the development of the document will be in the Introduction and/or on the ISO list of patent declarations received (see www.iso.org/patents).

Any trade name used in this document is information given for the convenience of users and does not constitute an endorsement.

For an explanation of the voluntary nature of standards, the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the World Trade Organization (WTO) principles in the Technical Barriers to Trade (TBT), see www.iso.org/iso/foreword.html.

This document was prepared by Technical Committee ISO/TC 209, *Cleanrooms and associated controlled environments*, in collaboration with the European Committee for Standardization (CEN) Technical Committee CEN/TC 243, *Cleanroom technology*, in accordance with the Agreement on technical cooperation between ISO and CEN (Vienna Agreement).

This second edition cancels and replaces the first edition (ISO 14644-9:2012), of which it constitutes a minor revision. The changes are as follows:

- "Class" (classification, classified) has been changed to grade or assessment where appropriate;
- ISO 14644-6 has been removed from the opening text of [Clause 3](#) and, as a result, [Clause 2](#);
- entry 3.8 removed from [Clause 3](#);
- ISO 4287 and ISO 4288 replaced by ISO 21920-2 and ISO 21920-3, respectively;
- ISO 16232-2, ISO 16232-3, ISO 16232-4 and ISO 16232-5 replaced by ISO 16232;
- minor editorial changes.

A list of all parts in the ISO 14644 series can be found on the ISO website.

Any feedback or questions on this document should be directed to the user's national standards body. A complete listing of these bodies can be found at www.iso.org/members.html.

Introduction

Cleanrooms and associated controlled environments provide for the control of contamination to levels appropriate for accomplishing contamination-sensitive activities. Products and processes that benefit from the control of contamination include those in such industries as aerospace, microelectronics, optics, nuclear and life sciences (pharmaceuticals, medical devices, food, healthcare).

ISO 14644-1 to ISO 14644-8, ISO 14698-1 and ISO 14698-2 deal exclusively with airborne particle and chemical contamination. Many factors, besides the assessment of surface cleanliness, should be considered in the design, specification, operation and control of cleanrooms and other controlled environments. These factors are covered in some detail in other parts of ISO 14644 and ISO 14698.

This document provides an analytical process for the determination and designation of surface cleanliness levels based on particle concentration. This document also lists some methods of testing, as well as procedure(s) for determining the concentration of particles on surfaces.

Where regulatory agencies impose supplementary guidelines or restrictions, appropriate adaptations of the testing procedures might be required.

NOTE When assessment of surface cleanliness by particle concentration (SCP) at critical control point(s) is used as an additional cleanliness attribute to classification of air cleanliness by airborne particle concentration in accordance with ISO 14644-1, then the space can be described as a cleanroom or clean-zone. If SCP is used alone, then the space is described as a controlled zone.

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Cleanrooms and associated controlled environments —

Part 9:

Assessment of surface cleanliness for particle concentration

1 Scope

This document establishes a procedure for the assessment of particle cleanliness levels on solid surfaces in cleanrooms and associated controlled environment applications. Recommendations on testing and measuring methods, as well as information about surface characteristics, are given in [Annexes A](#) to [D](#).

This document applies to all solid surfaces in cleanrooms and associated controlled environments, such as walls, ceilings, floors, working environments, tools, equipment and products. The procedure for the assessment of surface cleanliness by particle concentration (SCP) is limited to particles of between 0,05 µm and 500 µm.

The following issues are not considered in this document:

- requirements for the cleanliness and suitability of surfaces for specific processes;
- procedures for the cleaning of surfaces;
- material characteristics;
- references to interactive bonding forces or generation processes that are usually time-dependent and process-dependent;
- selection and use of statistical methods for assessment and testing;
- other characteristics of particles, such as electrostatic charge, ionic charges and microbiological state.

2 Normative references

There are no normative references in this document.

3 Terms and definitions

For the purposes of this document, the following terms and definitions apply.

ISO and IEC maintain terminology databases for use in standardization at the following addresses:

- ISO Online browsing platform: available at <https://www.iso.org/obp>
- IEC Electropedia: available at <https://www.electropedia.org/>

3.1
descriptor for specific particle size ranges
differential descriptor that expresses surface cleanliness by particle concentration (SCP) level within specific particle size ranges

Note 1 to entry: The descriptor may be applied to particle size ranges of special interest or those particle size ranges that are outside the range of the grading system and specified independently or as a supplement to the SCP levels.

3.2
direct measurement method
assessment of the contamination without any intermediate steps

3.3
indirect measurement method
assessment of the contamination with intermediate steps

3.4
solid surface
boundary between the solid and a second phase

3.5
surface particle
solid and/or liquid matter adhered and discretely distributed on a surface of interest, excluding film-like matter that covers the whole surface

Note 1 to entry: Surface particles are adhered via chemical and/or physical interactions.

3.6
surface cleanliness by particle concentration
SCP
condition of a surface with respect to its particle concentration

Note 1 to entry: The surface cleanliness depends upon material and design characteristics, stress loads (complexity of loads acting on a surface) and prevailing environmental conditions, along with other factors.

3.7
surface cleanliness by particle concentration level
SCP rating
grading number stating the maximum allowable surface concentration, in particles per square metre, for a considered size of particles [surface cleanliness by particle concentration (SCP) grades 1 to 8], where level 1 represents the cleanest level

3.8
surface particle concentration
number of individual particles per unit of surface area under consideration

4 Abbreviated terms

For the purposes of this document, the following abbreviated terms apply.

AFM	atomic force microscopy
CNC	condensation nucleus counter
EDX	energy dispersive X-ray spectroscopy
ESCA	electron spectroscopy for chemical analysis
ESD	electrostatic discharge

IR	infrared (absorption spectroscopy)
OPC	optical particle counter
PET	polyethylene terephthalate
SCP	surface cleanliness by particle concentration
SEM	scanning electron microscopy
UV	ultraviolet (spectroscopy)
WDX	wavelength-dispersive X-ray spectroscopy

5 The surface cleanliness level assessment system

5.1 ISO-SCP grading level format

The degree of SCP in a cleanroom or associated controlled environment shall be designated by a cleanliness level grading number, N , specifying the maximum total particle concentration on surfaces permitted for a considered particle size. N shall be determined from [Formula \(1\)](#) with the maximum permitted total particle concentration on the surface, $C_{SCP;D}$, in particles per square metre of surface, for each considered particle size, D :

$$C_{SCP;D} = k \frac{10^N}{D} \quad (1)$$

where

$C_{SCP;D}$ is the maximum permitted total surface concentration, in particles per square metre of surface, of particles that are equal to or larger than the considered particle size; $C_{SCP;D}$ is rounded to the nearest whole number, using no more than three significant figures;

N is the SCP cleanliness level grading number, which is limited to SCP grade level 1 to SCP grade level 8; the SCP grade level number N is qualified by the measured particle diameter D , in micrometres;

NOTE N refers to the exponent base 10 for the concentration of particles at the reference particle size of 1 μm .

D is the considered particle size, in micrometres;

k is a constant 1, in micrometres.

NOTE 1 The SCP grade level based on the particle concentration can be a time- and process-dependent value due to the dynamic characteristics of particle generation and transportation.

NOTE 2 Due to the complexity of statistical evaluations and readily available additional references, the selection and use of statistical methods for testing are not described in this document.

The concentration $C_{SCP;D}$, as derived from [Formula \(1\)](#), shall serve as the definitive value. [Table 1](#) presents selected SCP grading levels and corresponding maximum cumulative permitted total surface concentrations for considered particle sizes.

[Figure 1](#) provides a representation of the selected surface particle grade levels in graphical form.

Table 1 — Selected SCP grading levels for cleanrooms and associated controlled environments

Units in particles per square metre

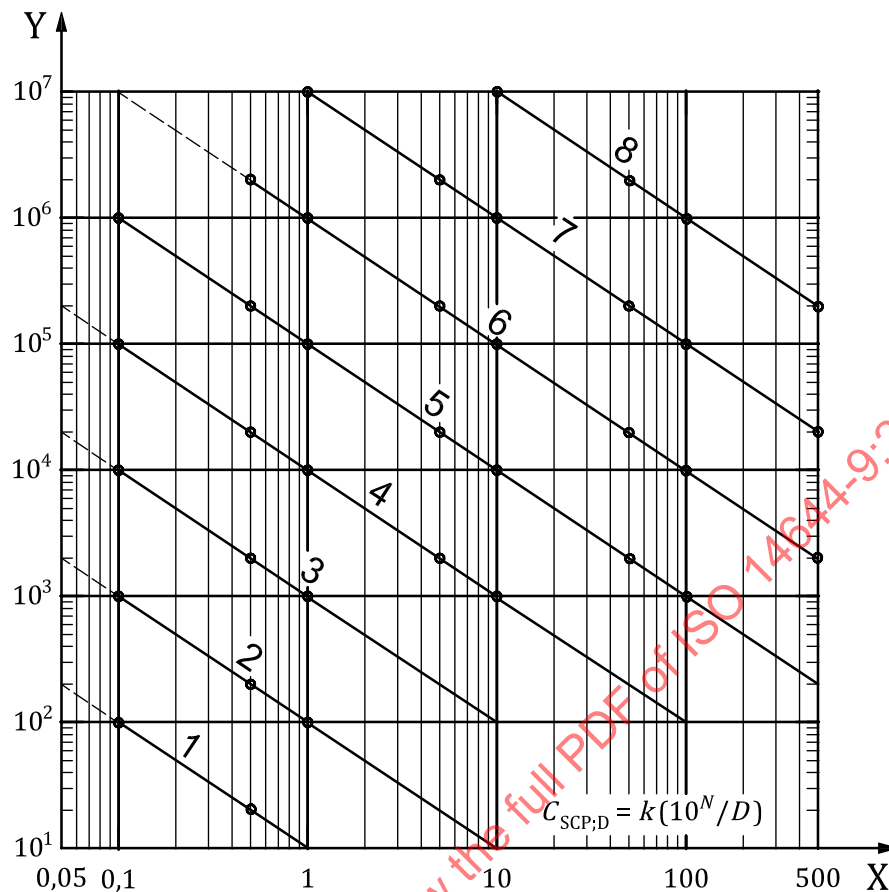
SCP level	Particle size								
	$\geq 0,05 \mu\text{m}$	$\geq 0,1 \mu\text{m}$	$\geq 0,5 \mu\text{m}$	$\geq 1 \mu\text{m}$	$\geq 5 \mu\text{m}$	$\geq 10 \mu\text{m}$	$\geq 50 \mu\text{m}$	$\geq 100 \mu\text{m}$	$\geq 500 \mu\text{m}$
SCP level 1	(200)	100	20	(10)					
SCP level 2	(2 000)	1 000	200	100	(20)	(10)			
SCP level 3	(20 000)	10 000	2 000	1 000	(200)	(100)			
SCP level 4	(200 000)	100 000	20 000	10 000	2 000	1 000	(200)	(100)	
SCP level 5		1 000 000	200 000	100 000	20 000	10 000	2 000	1 000	(200)
SCP level 6		(10 000 000)	2 000 000	1 000 000	200 000	100 000	20 000	10 000	2 000
SCP level 7				10 000 000	2 000 000	1 000 000	200 000	100 000	20 000
SCP level 8						10 000 000	2 000 000	1 000 000	200 000

The values in Table 1 are concentrations of particles of the related particle size and SCP level per surface area of one square metre (1 m^2) equal to or larger than the considered particle size ($C_{\text{SCP};D}$).

For figures in parentheses, the corresponding particle sizes should not be used for level determination purposes; select another particle size for more accurate determination.

The minimum area for testing should be statistically representative of the surface under consideration.

NOTE Assessment of the lower SCP levels requires numerous measurements to establish a significant value.



Key

- X considered particle size, D (μm)
- Y particle concentration on a surface $\geq D$, $C_{\text{SCP};D}$ (particles/ m^2)
- 1 SCP grade level 1
 - 2 SCP grade level 2
 - 3 SCP grade level 3
 - 4 SCP grade level 4
 - 5 SCP grade level 5
 - 6 SCP grade level 6
 - 7 SCP grade level 7
 - 8 SCP grade level 8

The solid lines shown on the graph shall be used for level assessment purposes. The dashed lines should not be used for level assessment purposes.

NOTE Particle distribution on surfaces typically is not a normal distribution, but is affected by different factors, such as roughness, porosity, electrostatic charge and deposition mechanisms (see [Annex A](#)).

EXAMPLE SCP grade level 5 ($1\ \mu\text{m}$) signifies that $1\ \text{m}^2$ of surface may carry a maximum of 10^5 particles with a considered particle size $\geq 1\ \mu\text{m}$ ($D = 1$). SCP grade level 5 ($10\ \mu\text{m}$) signifies that $1\ \text{m}^2$ of surface may carry a maximum of 10^4 particles per square metre with a considered particle size $\geq 10\ \mu\text{m}$ ($D = 10$). Any other measured particle size ($D = x$) which leads to a concentration that lies below the relevant SCP line is within the specification of SCP grade level 5 ($x\ \mu\text{m}$).

Figure 1 — SCP grade levels

For particle sizes outside the limits of the level numbering system and in cases where only a narrow particle range or individual particle sizes are of interest, a descriptor can be used (see [Annex B](#)).

5.2 Designation

The SCP grade level number shall be formatted as follows: SCP grade level N (D μm).

The designation of the SCP grade for cleanrooms and associated controlled environments shall also include the following:

- a) the surface type measured;
- b) the surface area measured;
- c) the measurement method applied.

Details of measurement methods applied, including sampling techniques and measurement devices, should be retrieved from test reports.

The considered particle size should be determined by agreement between the customer and supplier.

The SCP grade level shall be stated in relation to the measured particle size diameter.

EXAMPLE 1 SCP grade level 2 (0,1 μm); wafer or glass substrate, surface area: 310 cm^2 ; surface particle counter.

EXAMPLE 2 SCP grade level 5 (0,5 μm); inner wall of a bottle, surface area: 200 cm^2 ; liquid dispersion — liquid particle counter.

5.3 General information on surface cleanliness levels of particle concentration

Airborne particle concentration and surface particle concentration are generally related. The relationship is dependent on many factors, such as airflow turbulence, rate of deposition, time of deposition, deposition velocity, concentration within the air and surface characteristics such as electrostatic charge (see [A.2.4](#)).

To determine SCP, various parameters (see [Annex C](#)) and surface characteristics (see [Annex A](#)) that influence testing should be taken into account.

6 Demonstration of conformity

6.1 Principle

Conformity with SCP grade cleanliness level requirements, as specified by the customer, is verified by performing tests and by providing documentation of the results and conditions of the testing.

Details for demonstrating conformity (see [6.3](#)) shall be agreed upon between the customer and supplier in advance of testing.

6.2 Testing

Tests performed to demonstrate conformity shall be conducted in a controlled environment using suitable test methods and calibrated instruments, whenever possible.

Direct and indirect test methods can be used for demonstrating conformity and are given in [Annex D](#). The list of typical methods described is not exhaustive. Alternative methods of comparable accuracy may be specified by agreement.

NOTE Measurement by different methods, even when correctly applied, can produce different results of equal validity.

Repeated measurements are recommended.

The test method and environment shall be agreed upon between the customer and supplier.

Precautions should be taken to reduce electrostatic charge around the test zone, since electrostatic charge enhances particle deposition onto surfaces. If the surface is neither conductive, nor grounded or charge-neutralized, electrostatic charges can occur (see [Annex A](#)). Therefore, test results can vary.

6.3 Test report

The results from testing each surface shall be recorded and submitted as a comprehensive report, along with a statement of conformity or non-conformity with the specified SCP grade levels.

The test report shall include as a minimum the following:

- a) basic data:
 - date and time of testing;
 - name and address of the testing organization;
 - name of testing personnel;
- b) references consulted:
 - standards;
 - guidelines;
 - regulations;
 - number and year of publication of this document, i.e. ISO 14644-9:2022;
- c) environmental data:
 - environmental conditions for sampling (i.e. temperature, humidity, cleanliness);
 - environmental conditions for measurement (i.e. temperature, humidity, cleanliness) (not essential for use with direct methods);
 - location (e.g. room) used for the measurements;
- d) specimen:
 - clear identification of the test object;
 - description of the test object;
 - graph and/or sketch of the test specimen;
- e) test setup:
 - photo and/or sketch of the test setup;
 - description of operating parameters;
 - description of measurement points;
 - description of hardware used in the test setup;
- f) measurement devices:
 - identification of the instrument(s) and measuring devices used and current calibration certificate(s);
 - measurement range of measuring devices used;
 - reference of calibration certificates;

g) performing the test:

- relevant details of the test procedure used, with any available data describing deviations from the test procedure (if agreed);
- surface condition before sampling (e.g. after cleaning, after packaging, under atmospheric or vacuum conditions);
- specified test and measurement procedure or method;
- occupancy state(s) during sampling and measurement;
- specified test method(s);
- all agreed documentation (e.g. raw data, background particle concentrations, pictures, graphs, cleaning and packaging);
- duration, location and position of sampling (not essential for use with direct methods);
- duration, location and position of measurement (not essential for use with direct methods);
- noticeable observations made during sampling or measurement, where applicable;
- number of measurements performed;
- clear identification of the position and the area of the surface measured and specific designations for coordinates of the surface, if applicable;

h) results and analysis:

- visual inspection of the test surface before and after measurement, where applicable;
- measurement values and/or their analysis;
- statement of data quality;
- particle size ranges considered;
- test results, including particle concentration data for given particle sizes, for all tests performed;
- SCP grading level with designation expressed as SCP cleanliness grade level *N*;
- acceptance criteria for the clean surface, if agreed between the customer and the supplier.

Annex A (informative)

Surface characteristics

A.1 Surface description

A surface is commonly characterized by its texture (such as roughness, porosity), its mechanical properties (such as hardness) and its physicochemical properties (such as electrostatic surface charge and surface tension). Each of these properties should be considered before selecting a test method for the surface cleanliness assessment, or as an aid for the interpretation of the test results.

A.2 Surface characteristics

A.2.1 Roughness

A.2.1.1 Description

As the roughness of a surface affects many of its physical properties, surface roughness is not easily described by one single parameter, nor is it an intrinsic property of the surface. Roughness exists in two principal planes: at right angles to the surface, where it may be characterized by height, and in the plane of the surface, identified as “texture” and characterized by waviness. The roughness of a surface can be determined by mechanical or optical methods.

A.2.1.2 Testing

A frequently used mechanical method for the determination of roughness is the stylus instrument (see, for example, ISO 21920-2 or ISO 21920-3).

Frequently used optical methods for the determination of roughness and porous texture are microscopes (optical, confocal, interferometry, with or without tunnel effect, taper sectioning).

A.2.2 Porosity

A.2.2.1 Definition and description

Porosity is a measure of the void spaces in a material and is expressed as a decimal between 0 and 1, or as a percentage between 0 % and 100 %.

- **Effective porosity** (also called open porosity) refers to the fraction of the total volume in which fluid flow is effectively taking place (this excludes dead-end pores or non-connected cavities).
- **Macroporosity** refers to pores equal to or greater than 50 nm in diameter. Fluid flow through macropores is described by bulk diffusion.
- **Mesoporosity** refers to pores equal to or greater than 2 nm but less than 50 nm in diameter.
- **Microporosity** refers to pores smaller than 2 nm in diameter. Movement in micropores is by activated diffusion.

A.2.2.2 Testing

There are several ways to estimate the porosity of a given material or mixture of materials, which is called material matrix.

The **volume/density method** is fast and highly accurate (normally within $\pm 2\%$ of the actual porosity). The volume and the weight of the material are measured. The weight of the material divided by the density of the material gives the volume that the material takes up, minus the pore volume. Therefore, the pore volume is simply equal to the total volume minus the material volume, i.e. $(\text{pore volume}) = (\text{total volume}) - (\text{material volume})$.

The **water saturation method** is slightly more difficult but is more accurate and more direct. Take a known volume of the material and a known volume of water. Slowly dump the material into the water and allow it to saturate while pouring. Allow it to sit for a few hours to ensure that the material is fully saturated. Then remove the unsaturated water from the top of the beaker and measure its volume. The total volume of the water originally in the beaker minus the volume of water not saturated is the volume of the pore space, i.e. $(\text{pore volume}) = (\text{total volume of water}) - (\text{unsaturated water})$.

Mercury intrusion porosimetry requires the sample to be placed in a special filling device that allows the sample to be evacuated, followed by the introduction of liquid mercury. The size of the mercury envelope is then measured as a function of increased applied pressure. The greater the applied pressure, the smaller the pore entered by mercury. Typically, this method is used over the range of pores from 300 μm to 0,0 035 μm . Because of increased safety concerns over the use of mercury, several non-mercury intrusion techniques have been developed and should be considered as alternatives.

Nitrogen gas adsorption is used to determine fine porosity in materials. In very small pores, nitrogen gas condenses on pore walls that are less than 0,090 μm in diameter. This condensation is measured either by volume or weight.

A.2.3 Hardness

There are many National and International Standards on hardness tests for each material type. Hardness is frequently measured by the penetrating force of a diamond ball or tip, by the indentation of a hard body or by the rebound properties of an impactor.

The Rockwell, Brinell, Shore and Vickers method for metals is covered by ASTM E18-07. Geometry and pressure are chosen at the beginning of the test as a function of the thickness of the sample, the composition of the metal and the supposed hardness.

A.2.4 Static electricity

A.2.4.1 Definition and description

Static electricity is defined as an electrical charge caused by an imbalance of electrons on the surface of a material. This imbalance of electrons produces an electrostatic field that can influence the determination of the surface cleanliness of objects. ESD is defined as the transfer of charge between bodies at different electrical potentials.

Any relative motion and physical separation of materials or flow of solids, liquids or particle-laden gases can generate electrostatic charges. Common sources of ESD include personnel, items made from common polymeric materials and processing equipment. ESD can damage parts by direct contact with a charged source or by electric fields emanating from charged objects.

Charged surfaces can attract and hold particle contaminants. If the selected measurement method to determine the surface cleanliness is based on an indirect detection of particles on surfaces (see [D.2.3.3.5](#)), these measurement results might be inaccurate, as the particle removal is diminished. Therefore, especially when using indirect measurement methods, action should be taken to reduce ESD effects.

A.2.4.2 Testing

Determination of the ESD properties of the specimen surfaces can be helpful in estimating the influence of the removal efficiency of particles from surfaces (e.g. IEC 61340-5-1, ISO 10015, IEST RP-CC022.2, SEMI E43-0301, SEMI E78-0706).

A.2.5 Superficial tension

A.2.5.1 Definition

Superficial tension is the energy necessary to increase the surface by one area unit. It is usually defined as γ and expressed in joules per square metre (J/m^2) or in newtons per metre (N/m).

A.2.5.2 Testing

The best-known method is the measurement of the contact angle by the “set drop”, see Reference [19].

When a drop of liquid is brought into contact with a flat solid surface, its shape depends on the molecular force within the liquid for cohesive force, or between the liquid and solid for adhesive force. The contact angle between the liquid and solid is used as the surface tension index (see Figure A.1). It is generally found that liquids with low surface tension easily wet most solid surfaces, giving a zero-contact angle. The molecular adhesion between solid and liquid is greater than the cohesion between the molecules of the liquid.

The contact angle measurement is performed with an optical method ($\times 10$ to $\times 50$) magnifying the drop profile set on the flat solid surface.

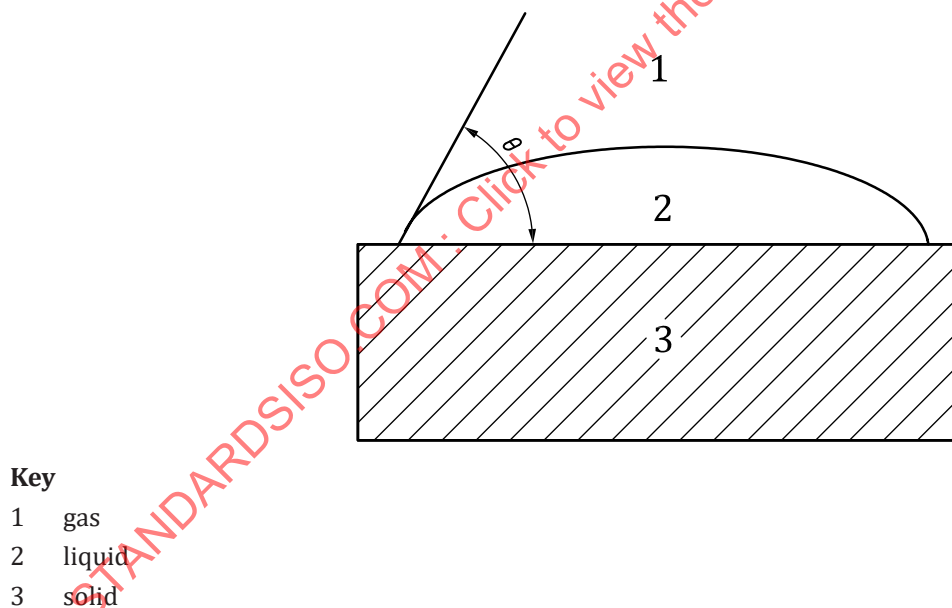


Figure A.1 — Shape of a drop of liquid in contact with a solid surface when the contact angle is $\theta < 90^\circ$

Annex B (informative)

Descriptor for specific particle size ranges

B.1 Application

For particle sizes outside the range of the cleanliness level grading system, a differential descriptor can be used. This descriptor can also be used for specific particle size ranges that are of special interest. In these cases, the descriptor can be used in addition to the SCP grade levels.

B.2 Surface descriptor for specific particle size ranges

The N_{ss} (particle number concentration of a specific particle size range) descriptor for specific particle size ranges may be specified independently or as a supplement to the SCP grading levels. The descriptor can be applied to any particle size range of special interest.

The surface particle concentration C_s within the particle size range D_L and D_U is a differential value.

The N_{ss} for a single particle size range is expressed according to [Formula \(B.1\)](#):

$$N_{ss}(C_s; D_L; D_U) \ a; \ b \quad (B.1)$$

where

C_s is the maximum permitted total surface concentration, in particles per square metre of surface, of the specified particle size range;

D_L is the lower limit of the specified particle size range, in micrometres;

D_U is the upper limit of the specified particle size range, in micrometres;

a is the measurement method used to determine particle size in the specified range;

b is the considered surface.

EXAMPLE 1 For the particle concentration on a metallic surface in the particle size range of 1 μm to 5 μm , the required value is 10 000 particles/ m^2 (1,0 particles/ cm^2). The particle concentration is measured by an optical microscope. The designation is:

$N_{ss}(10\ 000; 1; 5)$ optical microscope; metallic surface

If two or more size ranges are used, apply [Formula \(B.2\)](#). N_{ss} is then expressed in the format:

$$N_{ss} \begin{pmatrix} C_{s1}; & D_{L1}; & D_{U1} \\ C_{s2}; & D_{L2}; & D_{U2} \\ \dots & & \\ C_{si}; & D_{Li}; & D_{Ui} \\ \dots & & \end{pmatrix} \begin{pmatrix} a_1; & b \\ a_2; & b \\ \dots & \\ a_i; & b \\ \dots & \end{pmatrix} \quad (B.2)$$

where

- C_{si} is the maximum permitted total surface concentration, in particles per square metre of surface, of the i -th particle size range;
- D_{Li} is the lower limit of the i -th particle size range, in micrometres;
- D_{Ui} is the upper limit of the i -th particle size range, in micrometres;
- a_i is the measurement method used to determine particle size in the i -th range;
- b is the considered surface.

EXAMPLE 2 For the particle concentration on a glass plate, based on the simultaneous use of a scattered-light scanner in the particle size range of 0,1 µm to 0,5 µm, and an optical microscope in the particle size range of 5 µm to 20 µm, the measured values are 9 000 particles/m² (0,9 particles/cm²) and 500 particles/m² (0,05 particles/cm²), respectively. The values are within the maximum allowable limits of 10 000 particles/m² and 500 particles/m², respectively. The designation is:

$$N_{ss} \left(\begin{array}{ccc} 10\,000; & 0,1; & 0,5 \\ 500; & 5; & 20 \end{array} \right) \begin{array}{l} \text{scattered light scanner; glass plate} \\ \text{optical microscope; glass plate} \end{array}$$

When measurement methods and/or specific surfaces are not predefined or are not essential, designations a and b can be omitted. In this case, the descriptor is expressed according to [Formula \(B.3\)](#):

$$N_{ss} (C_s; D_L; D_U) \quad (B.3)$$

where

- C_s is the maximum permitted total surface concentration, in particles per square metre of surface, of the specified particle size range;
- D_L is the lower limit of the specified particle size range, in micrometres;
- D_U is the upper limit of the specified particle size range, in micrometres.

Where only one particle size is of interest, the lower and upper limit in [Formula \(B.3\)](#) can be used to frame the particle size of interest through an agreement between the customer and supplier.

EXAMPLE 3 For the particle size 5 µm, the required value is 200 particles/m² (0,02 particles/cm²). D_L can be set to 4,5 µm and D_U can be set to 5,5 µm. The designation is:

$$N_{ss} (200; 4,5; 5,5)$$

When measurement methods and/or specific surfaces are not predefined or are not essential, designations a and b can be omitted. The descriptor for two or more particle size ranges is expressed according to [Formula \(B.4\)](#):

$$N_{ss} \left(\begin{array}{ccc} C_{s1} ; & D_{L1} ; & D_{U1} \\ C_{s2} ; & D_{L2} ; & D_{U2} \\ \dots & & \\ C_{si} ; & D_{Li} ; & D_{Ui} \\ \dots & & \end{array} \right) \quad (B.4)$$

EXAMPLE 4 For the particle concentration in the particle size ranges of 0,1 µm to 0,5 µm and 5 µm to 20 µm, the measured values are 9 000 particles/m² (0,9 particles/cm²) and 500 particles/m² (0,05 particles/cm²), respectively. They are within the maximum allowable limits of 10 000 particles/m² and 500 particles/m², respectively. The designation is:

$$N_{ss} \begin{pmatrix} 10\,000; & 0,1; & 0,5 \\ 500; & 5; & 20 \end{pmatrix}$$

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Annex C (informative)

Parameters influencing the SCP grading level assessments

C.1 Background

Parameters that can influence the testing and measuring of surfaces are presented in [C.2](#). The information is not exhaustive and there is no ranking. More detailed information on measurement methods and surface characteristics is given in [Annex D](#).

C.2 Parameters

C.2.1 Physical or chemical properties

- **Surface energy states.** The attraction and removal of particles might be influenced, for example, by cohesive or adhesive characteristics and/or hydrophilic or hydrophobic properties of the surface.
- **Porosity of the surface.** In most cases, the higher the degree of porosity, the more complex the distinction between surface imperfection and particle detection.
- **Cleanability of the surface.** When surfaces are difficult to clean, the discrimination between surface imperfection and particle detection is complex.
- **Optical characteristics of the surface.** When direct test methods are applied, different optical characteristics of the surface to be tested will lead to different measurement results. This difference is not noted for indirect methods.
- **Electrostatic properties of the surface.** The electrostatic properties of the surface will influence the attraction and removal of ESD-charged contamination.
- **Magnetic characteristics of the surface.** The magnetic characteristics of the surface will influence the attraction and removal of materials with magnetic properties.

C.2.2 Shape of the surface and particles

- **Morphology of the particle** (e.g. round, flat, oval, peaked) and topography of the surface can influence the measurement result.
- **Surface condition** (e.g. being cleaned, extruded, polished). The particle attachment force varies depending on the surface condition. This variation also affects the removal efficiency of particles and the ability to distinguish between particles, roughness and porosity.
- **Roughness, porosity or waviness of the surface.** The roughness, porosity or waviness will have an effect on the efficiency of removing particles for indirect methods.
- **Shape or geometry of the particle.** The shape of particles can also influence the measurement results. For example, long particles and perfectly round particles can lead to the same optical particle counter values after detaching the particles from the surface, but to completely different gravimetric values. In general, fibres are considered as particles having an aspect (length-to-width) ratio of 10 or more.

C.2.3 Ability to measure or analyse and appropriate statistics for particle analysis

The number of measurements should lead to statistically significant results. Therefore, an appropriate statistical method with regard to the frequency distribution of individual measurements should be used to estimate the confidence limit.

- **Ability to measure or analyse.** Feasibility to perform the measurements, depending on the accessibility of a measurement device to the specimen.

EXAMPLE Performing measurements within holes or microtubes.

- **Ability to detect particles** (direct or indirect methods). The ability to distinguish between surface sedimented particles and surface imperfections (direct method) or the detachability of surface sedimented particles (indirect method) will influence the SCP grade level.
- **Geometric size of surface area to be measured.** Depending on the geometric size of the surface area to be measured, different methods should be selected. In the majority of cases, the statistics for the number of samples to be taken and how to analyse the measurement values should be developed individually.
- **Inline capability.** Several measurements should be taken for statistical significance. The influence of measurement repeatability is diminished with inline options.
- **Frequency distribution of individual measurements.**
- **Particle size(s) to be measured.**
- **Distribution of particles on surface area.**

C.2.4 Particle origin

Particles on the surface can originate from, for example, material friction, deterioration, deposition of airborne particles or chemical reaction of gaseous matters with the surface, forming solid or liquid products.

EXAMPLE $2\text{NH}_3 + \text{H}_2\text{O} + \text{SO}_3 \rightarrow (\text{NH}_4)_2\text{SO}_4$.

Annex D (informative)

Measurement methods for determining surface cleanliness by particle concentration

D.1 Surface cleanliness by particle concentration

In order to obtain quantitative information on surface cleanliness, appropriate measurement methods should be selected. In some cases where quantitative information cannot be ascertained on a surface, it is possible at least to obtain a qualitative result. Qualitative results cannot be used for the SCP grades defined in [Clause 5](#).

D.2 Criteria for the measurement of surface cleanliness by particle concentration

D.2.1 General

The cleanliness of surfaces can be assessed as soon as the particle contamination is detectable.

As a quantitative criterion for the evaluation and grading of the surface cleanliness, the number of all adherent particles of undesired material should be determined. It should be possible to determine the dimension and number in relation to the contaminated surface area (surface particle contamination).

NOTE The cleanliness assessment of textiles and/or porous surfaces also takes into account particles which might emanate from the specimen.

D.2.2 Requirements of the measurement method

The measurement method is essentially selected according to criteria determined by the surface being tested and its characteristics. Some of the most important requirements are summarized as follows:

- information regarding particle characteristics (e.g. particle size, concentration, size distribution, material, shape, position);
- feasible measuring positions (transportable measuring device, i.e. applicable even to large immovable surfaces);
- measurements independent of surface characteristics (e.g. roughness, waviness, component shape);
- testing speed and effort involved (i.e. use of random sampling or series testing);
- flexibility (i.e. whether the method can be rapidly implemented on various surfaces of different components);
- little or no surface alteration caused by the measurement procedure (i.e. measurement surfaces are not altered as a result of being wetted with flushing fluids).

Due to the requirements mentioned, the measurement methods described in [D.2.3](#) can be listed and limited for each of the applications.

D.2.3 Measurement methods

D.2.3.1 General

Ideally, surface cleanliness is assessed when test surfaces are of a low roughness and can be accessed by the chosen measurement device. In principle, the following methods can be utilized to measure the particle cleanliness of surfaces:

- direct methods;
- indirect methods.

In most cases, direct methods that do not require sample taking should be given priority. Generally, these methods involve less measurement activity and are associated with fewer errors, thus giving more reproducible results than indirect methods. However, depending on component and manufacturing requirements (complex component shape, rough surface), indirect methods are often the only possible alternative in order to determine the particle cleanliness of a surface.

D.2.3.2 Direct methods

Particles are recorded and measured directly on the test surface. Neither the component surface nor the particles present on it should be altered or affected by the measurement. If a testing surface requires transport to a measuring device, the transportation (handling, packaging) should be carried out so as to avoid any further contamination from reaching the surface.

The methods are given in [Table D.1](#) and have been characterized here according to the requirements to be fulfilled and the limitations of the methods.

Table D.1 — Comparison of measurement methods for the direct detection of particles on surfaces

Method ^a	Detection limits	Determination of concentration	Size distribution	Material analysis	Shape analysis	Determination of position	Transportability	Independent of surface	Accessibility	Testing speed	Flexibility	Influence on surface
Visual inspection	> 25 µm	+	+	+	+	+	++	+	+	++	++	++
Light microscope (with image-processing)	> 4,0 µm	++	++	+	++	++	+	+	+	++	++	++
Oblique-, glancing-, side-light systems (with image-processing)	> 0,5 µm	++	++	+	+	++	++	+	+	++	++	++
Scattered-light scanner	> 0,07 µm	++	++	—	++	++	—	—	—	++	—	++
SEM	> 0,01 µm	+	+	++	++	++	—	—	—	—	+	+
AFM	> 0,01 µm	+	+	++	+	++	+	+	+	—	+	+
Key + + highly suitable + partially suitable — unsuitable or not useful ^a Particles that are not perfectly round should be measured on their largest axis.												

The efficiency of optical methods to detect particles on surfaces can be enhanced by the additional use of suitable light sources that are based on specific material effects (e.g. UV-, IR-activity). If, for instance, a UV-lamp is used, particles that are physically UV-active will be detected with a much better contrast.

Within the visible light spectrum, the colour/reflection grade of particles can be used for further discrimination of particle contamination.

D.2.3.3 Indirect methods

D.2.3.3.1 General

Since direct counting is often not possible on the surface of interest due to either object-related or measurement-related reasons, samples should be prepared before inspection. The particles to be recorded are detached from the testing surface (sampling) and are placed on or in a replacement substrate or medium. The particles are then measured using a technique adapted to the replacement substrate or medium concerned (see IEST-STD-CC1246D).

The measuring efficiency is reduced if not all particles that settled on surfaces can be detached from the testing surface by indirect methods.

Due to physical or chemical effects, such as adhesion or cohesion, or electrostatic forces, the detachment force can be insufficient, resulting in diminished measuring efficiency of surface particle contamination. Therefore, direct measurement is preferred.

However, care should be taken if more aggressive methods are used to remove particles from a surface, as such methods can erode a surface and form additional particles. The test method should be evaluated for this potential effect.

When using indirect methods, the background particle contamination of the intermediate media (e.g. flushing medium) should be known.

D.2.3.3.2 Detachment techniques

In the case of surfaces that are difficult to access due to the complex shape of a component, drawing samples with detached particles is often the only way of assessing SCP. The smaller the particles become, the more difficult it will be to detach them from surfaces, as several surface forces (e.g. electrostatic, cohesion, adhesion, capillary) increase. For the detachment of particles from tested surfaces, techniques such as the following can be used:

- **Tape-lifting.** The particles to be recorded are detached using a clean adhesive replacement substrate (adhesive tape or stamp) (see ASTM E1216-06) and guided directly to a measurement process (see ASTM F312-08).
- **Flushing.** The particles to be recorded are rinsed off using a clean flushing medium, such as gas or liquid (see ASTM F24-09). The particles contained in the measurement media are then examined using appropriate measuring devices (e.g. optical particle counters for gases) or the particles held in the medium are deposited on a replacement substrate (filtration, impaction) and then measured.

In all sampling cases, care should be taken to ensure that samples are not further contaminated by the devices or media used or by personnel, as this contamination affects the measurement result. The contamination brought in from the environment, processes and materials used for the inspection procedure should be less than 10 % of the presumed or specified particle numbers, at the relevant sizes, with each number calculated being rounded down (see ISO 16232). Furthermore, the sampling method chosen should be capable of detaching the required particles reliably and completely from the testing surface. Flushing the sampling equipment should be capable of ensuring the complete transfer of the particles present to the measurement medium or replacement surface. To optimize the sampling method, blank samples (i.e. sampling on a clean surface) or samples from surfaces contaminated in a defined way can be used.

D.2.3.3.3 Measurement of the medium being used for detaching surface sedimented particles

The particles present in the gaseous or liquid flushing medium that was used for detaching surface sedimented particles can be measured directly using appropriate optical particle counters (see [D.2.7.4](#)). The low test-flow rate of optical particle counters does not permit large volumes of flushing media to be measured completely. Therefore, a representative sample should be taken from the flushing volume. Especially in the case of large particles ($>3\ \mu\text{m}$), care should be taken to avoid any separating out or sedimentation of the particles as this would create an inaccurate measurement result. As with sampling techniques, sampling equipment and associated tubing should be kept clean when measuring flushing media.

D.2.3.3.4 Collection method

The particles detached from the testing surface are present in either a gaseous or a liquid flushing medium. In order to measure the number of the particles, they are first deposited on a surface. The substitute surface is then passed to the corresponding measurement systems for examination. For particle collection, methods and devices such as the following can be used:

- **Filtration systems** (liquid drawn through a filter membrane via a sucking strainer or liquid poured onto a filter medium). The flushing medium is guided through filter membranes possessing an appropriate pore size for the particle size to be determined. The filters containing the particles are then dried and analysed either gravimetrically (see [D.2.7.7](#)) or microscopically (see [D.2.6.2](#) and [D.2.7.2](#)).
- **Impactors**. Particles are deposited from gaseous flushing media onto an impaction plate. The impaction plates are then analysed microscopically.

All devices and handling steps associated with the collection process are subject to correspondingly high cleanliness requirements. Their level of cleanliness should be stated using blank samples.

D.2.3.3.5 Most frequently used indirect methods (see Table D.2)

For detachable particle size or efficiency of indirect measurement methods, smaller particles (smaller than approximately $1\ \mu\text{m}$) require a greater effort for removal from the surface for indirect measurement. The efficiency of a given method in detaching particles from surfaces is not only dependent on the particle size, but also related to the following parameters:

- shape and material of particles;
- existence of surface forces (e.g. electrostatic, cohesion, adhesion, capillary);
- detaching method (e.g. ultrasonic, megasonic, flushing, purging, blowing, drawing).

Due to the different methods used to overcome forces between particles and surfaces, and the interaction between these factors, the efficiency to detach particles varies significantly. Therefore, discrete values for the efficiency of indirect methods cannot be given.

To obtain additional information from indirect methods, analytical methods such as ESCA, EDX, Raman, UV or IR spectroscopy can be used for the characterization of particles.

Table D.2 — Comparison of measurement methods for the indirect detection of particles on surfaces

Method ^a	Limit of the detachment procedure	Estimation on limits of the measurement method	Determination of concentration	Size distribution	Material analysis	Shape analysis	Determination of position	Transportability	Independent of surface	Accessibility	Testing speed	Flexibility	Influence on surface
Examination of flushing medium (liquid/gaseous) using extinction particle counters (>1 µm)	0,2 µm	> 1 µm	++	++	—	—	—	++	++	++	+	++	—
Filtration or impaction of flushing medium and microscopic analysis (>0,5 µm)	0,2 µm	> 1 µm	++	++	+	++	—	+	++	++	—	++	—
Examination of flushing liquid using an OPC (start with flushing particles off the surface then draw them through an OPC) (>0,05 µm)	0,2 µm	> 0,2 µm	++	++	—	—	—	++	++	++	+	++	—
Examination of gaseous medium using an OPC (start blowing particles off the surface then draw them through an OPC) (>0,05 µm)	0,3 µm	> 0,3 µm	++	++	—	—	—	++	++	++	+	++	—
Filtration of flushing medium and gravimetric analysis (>0,1 mg)			++	—	—	—	—	++	++	++	—	++	—
Key + + highly suitable + partially suitable — unsuitable or not useful NOTE Use of microscopic or gravimetric analysis: whether filters can be examined with microscopic or gravimetric analysis depends on the total number of particles found on the filters. The size of the particles is not a decisive factor. Empirically determined reference value: microscopic analysis is not possible for contamination greater than 3 mg on the surface of a filter (standard filter size of 47 mm) (see ISO 16232). Gravimetric analysis is not suitable for assessment of SCP levels, as no discrete single particles will be measured. Gravimetric analysis is used to determine the total mass of all contamination detached from the test surfaces. ^a Figures in parentheses are detection limits of the measurement devices.													

D.2.4 Determination of number of samples

The number of measurement points used and the overall surface under investigation determine the statistical certainty of the measurement results. As the measurement results in general are dependent