
**Respiratory protective devices — Terms,
definitions, graphical symbols and units
of measurement**

*Appareils de protection respiratoire — Termes, définitions, symboles
graphiques et unités de mesure*

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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 16972 was prepared by Technical Committee ISO/TC 94, *Personal safety — Protective clothing and equipment*, Subcommittee SC 15, *Respiratory protective devices*.

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Respiratory protective devices — Terms, definitions, graphical symbols and units of measurement

1 Scope

This International Standard is applicable to respiratory protective devices. It defines commonly used terms and specifies units of measurement to achieve a uniform interpretation and to prevent ambiguous use. It indicates graphical symbols that may be required to be placed on respiratory protective devices (RPD) or parts of RPD or instruction manuals, in order to instruct the person(s) using the RPD about its operation.

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 7000:2004, *Graphical symbols for use on equipment — Index and synopsis*

IEC 80416-1¹⁾, *Basic principles for graphical symbols for use on equipment — Part 1: Creation of graphical symbols for registration*

ISO 80416-2²⁾, *Basic principles for graphical symbols for use on equipment — Part 2: Form and use of arrows*

3 Terms and definitions

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

NOTE The terms are listed in alphabetical order. Terms which refer to current RPD are not listed in these definitions because they are design related and are described in Annex A. Bold type used within a definition identifies terms defined elsewhere in this list of preferred terms.

3.1

abrasion resistance

ability of an **RPD** and/or its components to withstand degradation from abrasive effects (e.g. scratch, scrape, scuff)

3.2

abrasive blasting respiratory protective device

device designed to protect the wearer from inhalation of, impact of, and abrasion by materials used or generated in abrasive blasting

3.3

accessory

item, or items, that are attached to the **RPD** that are not necessary for the RPD to meet the requirements of the ISO RPD performance standard and do not compromise its protection

3.4

adequacy assessment

selection method identifying **RPD** able to reduce the wearer's hazard exposure to acceptable levels

3.5

aerodynamic diameter

diameter of a unit density sphere having the same settling velocity as the particle in question

3.6

aerosol

suspension of solid, liquid or solid and liquid particles in a gaseous medium

3.7

aerosol penetration

ability of particles to pass through a particle filtering material

3.8

air flow resistance

pressure difference between upstream and downstream locations caused by the flow of air through parts and components of RPD such as, an exhalation valve, inhalation valve, filter(s), and tube, etc.

1) Cancels and replaces ISO 3461-1:1988.

2) Cancels and replaces ISO 4196:1984.

3.9

air-purifying respiratory protective device

device in which ambient air is passed through an air-purifying element(s) that remove(s) the contaminant(s)

NOTE Air is passed through the air-purifying element by means of the breathing action or by a blower.

3.10

ambient air bypass

means to enable the wearer to breathe the **ambient atmosphere** before entering and after leaving a hazardous atmosphere

3.11

ambient air system

device used to deliver ambient air at a low pressure directly to an atmosphere supplying **RPD** (manually or power assisted)

3.12

ambient atmosphere

air surrounding the **RPD** wearer

3.13

ambient concentration

concentration of a compound in the air surrounding the **RPD** wearer

3.14

ambient laboratory conditions

atmosphere where the temperature is between 16 °C and 32 °C and the relative humidity is between 20 % and 80 %

3.15

atmospheric contaminant

potentially harmful substance that is present in unacceptably high concentration in the ambient air, and to which workers may be exposed in their working environment

3.16

atmospheric dew point

temperature at which moisture begins to condense from a gas as the gas is cooled at standard atmospheric pressure. (1 013,25 hPa)

3.17

biological monitoring

analysis of exhaled air, a biological fluid (e g., urine, blood, perspiration), or a body tissue (e g., hair, nails) to assess the extent to which an individual has been exposed to a contaminant/hazard

3.18

body harness

means to enable a wearer to wear certain components of an **RPD** on the body

3.19

breakthrough concentration

concentration of test gas in effluent air at which a **gas filter** undergoing a gas filter capacity test is deemed exhausted

3.20

breakthrough curve

graph for gas filter(s) showing the relationship between effluent concentration of the test substance versus time of **gas filter(s)**

3.21

breakthrough time

time from the start of a **gas filter** capacity test to the time when the **breakthrough concentration** has been reached in effluent air under the defined conditions

3.22

breakthrough time curve

graph showing the relationship between **test concentration** and **breakthrough time** for **gas filter(s)**

3.23

breathable gas

mixtures of gases that are suitable for respiration without adverse effects to health

3.24

breathable gas cylinder

integral part of the **RPD** that contains the **breathable gas** supply

3.25

breathable gas quality

composition of a breathable gas as defined in relevant standards

3.26

breath cycle

time period comprised of one inhalation and one exhalation sequence

3.27

breathing bag

component of an **RPD** which compensates for variations in the **breathable gas** supply or demand and provides for peak inhalation flow requirements

3.28**breathing gas regeneration**

process whereby an RPD absorbs carbon dioxide in exhaled gas and delivers oxygen, controls water vapour and temperature of gas to be rebreathed

3.29**breathing machine**

ventilation machine that simulates respiratory ventilation by using waveforms, which can be sinusoidal or representative of the inhalation and exhalation cycle

NOTE

See also **metabolic simulator**.**3.30****breathing peak resistance**

maximum differential pressure of an RPD during inhalation (inhalation peak resistance) or exhalation (exhalation peak resistance)

3.31**breathing resistance**

pressure differential between upstream and downstream location caused by an RPD to the flow of breathable gas during inhalation (inhalation resistance) or exhalation (exhalation resistance)

3.32**BTPS**

body temperature pressure saturated standard condition for the expression of ventilation parameters

EXAMPLE Body temperature (37 °C), atmospheric pressure (1 013,25 hPa) and saturated air (vapour pressure = 62,66 hPa).

3.33**buddy breathing**

practice that enables a second person to simultaneously share the same **breathable gas** supply as that of the wearer of such RPD while both persons are attempting to move to a safe location

3.34**bypass valve**

component part of an RPD, which is furnished as an emergency manual valve to supply necessary **breathable gas** when the ordinary supply path is out of order

3.35**CO₂ concentration limits**

maximum allowed concentration of carbon dioxide within inhaled **breathable gas**

3.36**char length**

length of brittle residue found when a fabric or material is exposed to thermal energy

3.37**checking device**

characteristic of RPD to enable the wearer to verify, before use or periodically during use, that the manufacturer's minimum design conditions are met

3.38**cleaning/disinfection resistance**

ability of the device to withstand cleaning and disinfection processes defined by the manufacturer

3.39**clogging**

accumulation of particles on a filter with a consequent increase in its resistance to flow

3.40**combination filter**

filter intended to remove dispersed solid and/or liquid particles and specified gases and vapours from the flow of air passing through it

3.41**combined RPD**

RPD which are capable to operate in either filtering or breathable gas supply mode

3.42**communication performance**

ability of wearer to communicate face to face with others at a given distance

3.43**compatibility**

ability of RPD to be used in conjunction with other PPE

3.44**competent person**

person with suitable and sufficient experience and with practical and theoretical knowledge of the elements of an RPD programme for which (s)he is responsible

3.45**confined space**

area with limited access, as described in national regulations, which requires special considerations for entry

3.46

continuous flow valve

control valve which provides the wearer of a breathable gas-supplying **RPD** with **breathable gas** and allows the wearer to regulate a continuous air flow within prescribed limits

3.47

count median diameter

CMD

particle size of a particle distribution for which one-half the total number of particles are larger and one-half are smaller

3.48

dead space

space in which exhaled gas has not been purged and is subject to being rebreathed

3.49

demountable parts

parts of the device that are designed to be disconnected and reconnected by hand, without using tools

3.50

design duration

length of time specified by the manufacturer for a device in which the design conditions (e.g. flow rate) are met

3.51

dew point

temperature of air at a specified pressure when condensation of water will occur

3.52

doffing

process of removing or taking off the **RPD** effectively

3.53

donning

process of putting on the **RPD** effectively

3.54

drip

to run or fall in drops or blobs

3.55

dummy head

fixture simulating human heads, used in testing **RPD**.

3.56

durability

capability of the device to withstand all usage stress factors without degradation of performance or integrity

3.57

dust

aerosol consisting of mechanically produced solid particles derived from the breaking up of larger particles ranging from submicroscopic to macroscopic

NOTE

See also **fume** and **smoke**.

3.58

emergency breathing component

component of the **RPD** that comes into operation when the normally operating mode of the device is not functioning

NOTE

It provides an adequate level of protection for a period to enable the device wearer to exit the work area, unassisted, to a place of safety.

3.59

electromagnetic compatibility

EMC

ability of a device, unit of equipment or system to function satisfactorily in its electromagnetic environment without introducing intolerable electromagnetic disturbances to anything in that environment

3.60

end of battery life

lowest capacity level of a battery which still allows a proper continuous function of the **RPD**

3.61

end of life cycle indicator

system that indicates the end of the overall life of an **RPD**

3.62

end-of-service-life indicator

ESLI

system that warns the wearer of a **gas filtering RPD** of the approach of the end of adequate respiratory protection

3.63

environment

potential hazardous area in which the **RPD** is to be used

3.64

ergonomic factors/parameters

all aspects of the **RPD** that are intended to maximize the wearer's ability to perform their tasks while minimizing wearer fatigue and discomfort

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3.65**escape RPD**

respiratory protective device to be used only when escaping (emergency egress) from a dangerous **environment**

3.66**exhalation valve**

non-return valve which allows the release of exhaled and excess **breathable gas** from the **RPD**

3.67**explosive atmosphere**

mixtures of substances with air, under atmospheric conditions, in the form of gases, vapours, mists or dusts in which, after ignition has occurred, combustion spreads to the entire unburned mixture

3.68**exposed parts**

any part of the **RPD** which may come in contact with the ambient atmosphere during foreseeable conditions of use

3.69**eye irritation (external)**

redness, itching or tearing of the eyes caused by gases, vapours, particles and liquids

3.70**eye irritation (internal)**

redness, itching or tearing of the eyes caused by the **RPD** itself

3.71**fabric component**

any single, or combination of, natural or synthetic material(s) that are pliable and that are made by weaving, felting, forming, or knitting

3.72**face seal leakage**

leakage between the face and the **respiratory interface**

3.73**field of vision**

area of sight while wearing an **RPD**

3.74**filter holder**

component which is attached to either a respiratory interface or other part of the device and into which a filter, either encapsulated or unencapsulated, is inserted

3.75**filtering device**

RPD in which air passes through filter(s) before being inhaled

NOTE The device can be unassisted, power assisted or powered.

3.76**fit test**

use of a challenge agent and specific protocol to qualitatively or quantitatively determine an individual's ability to obtain an adequate seal with a specific make, model, and size of an **RPD**

3.77**flow rate**

air supply volume (mass) of **breathable gas** passing through the device in a given time

3.78**fogging**

reduction of the field of vision and visual acuity caused by condensation of humidity inside the **visor**

3.79**fume**

solid aerosols formed when the material from a volatilised solid condenses in air

3.80**gas**

fluid that is in a gaseous state at **standard temperature and pressure** that expands to occupy the space or enclosure in which it is confined

3.81**gas filter**

filter intended to remove specific gases and vapours from the atmosphere passing through it

3.82**gas filter capacity**

volume or mass of a gaseous contaminant that a **gas filter** can adsorb, absorb or catalyze the breakdown of

3.83**gas filter change schedule**

time interval after which a used **gas filter** is replaced with a new one

3.84**gas filtering RPD**

device consisting of a **respiratory interface** with a filter which removes certain gases or vapours from the air to be inhaled by the **wearer** for a limited period

3.85

hazardous atmosphere

any atmosphere that is **oxygen-deficient**, exceeds an **occupational exposure limit**, presents a fire/explosion hazard, and/or contains an airborne toxic or disease-producing contaminant in concentrations deemed to be hazardous

3.86

hazard ratio

estimated/measured airborne concentration of a substance divided by the **occupational exposure limit**

NOTE This ratio is calculated for each gas, vapour, and/or particulate component, or for a mixture when the components have additive effects, that pose a respiratory hazard.

3.87

head harness

means of holding a **respiratory interface** in place on the head

3.88

heads up display

HUD

visual monitor or warning in the line of sight of the **RPD** wearer

3.89

high pressure

equal to or greater than 1 MPa (10 000 mbar) absolute pressure

3.90

hose

hollow flexible conduit to carry **breathable gas** designated as **low**, **medium** or **high pressure**

3.91

hydration facility

design feature of **RPD** to allow the **wearer** to drink without **doffing** of **RPD**

3.92

hydrostatic test

calibrated expansion pressure test of the structural integrity of cylinders

3.93

hypoxia

any condition in which there is an inadequate supply of oxygen in the tissues

3.94

immediately dangerous to life or health

IDLH

atmosphere that poses an immediate threat to life, would cause irreversible adverse health effects, or would impair an individual's ability to escape from a dangerous atmosphere

3.95

impact resistance

ability of **RPD** to withstand mechanical shock and dynamic stress from environment

3.96

inhalation valve

valve which opens during inhalation and closes during exhalation

3.97

inhaled air

air breathed in by the wearer

3.98

integral filter

filter which is not separable from the rest of the **respiratory interface**

3.99

interactive flow rate

actual **RPD** breathing gas flow per time as a result of the wearer's demand

3.100

intrinsic safety

various designations that ensure the **RPD** is not a source of ignition in explosive atmospheres

3.101

intrinsically safe (IS) RPD

respiratory protective device that has been certified as not providing a source of ignition in an explosive atmosphere

3.102

inward leakage

leakage of the ambient atmosphere into the **respiratory interface** from all sources excluding filter(s) where present, when measured in the laboratory in the specific test atmosphere

NOTE It is expressed as a ratio of contaminant concentration inside an **RPD** and ambient atmosphere, as follows:

$$\text{Inward leakage (\%)} = C_i/C_o \times 100$$

where

C_i is the concentration of challenge agent inside the respiratory interface;

C_o is the concentration of challenge agent outside inside the respiratory interface.

3.103**laboratory gas filter capacity**

mass/volume of a specific test agent that will be removed or retained by certain mechanisms (e.g. physisorption, chemisorption, catalytical conversion, etc.) under specified conditions of temperature, relative humidity, challenge test gas concentration and the flow rate

NOTE The mass/volume is determined by measuring the breakthrough time at a defined breakthrough concentration.

3.104**leak-tightness**

ability to withstand a loss of pressure inside an RPD over a given time as determined by a laboratory test

3.105**life cycle**

time between date of manufacturing of the device to the date when device has to be withdrawn from service

3.106**liquefied-breathing gas**

oxygen or air stored in liquid form and supplied to the wearer in a gaseous form

3.107**loose fitting respiratory interface**

respiratory interface which does not rely on forming a complete seal to the wearer's skin

3.108**low boiling organic compound**

organic compound having a boiling point $\leq 65\text{ }^{\circ}\text{C}$ at atmospheric pressure

3.109**low pressure**

pressure up to 100 hPa (mbar) above atmospheric pressure (gauge)

3.110**manufacturer's minimum design condition**

lowest level of operating condition of the device as stated by the manufacturer at which the complete device will still meet the requirements for the designated class

3.111**manufacturer's minimum design flow rate****MMDF**

minimum air flow rate, as stated by the manufacturer, at which the relevant requirements are met

3.112**marking**

information included on the device to indicate specific RPD characteristics

3.113**mass median aerodynamic diameter****MMAD**

point in an aerodynamic particle size distribution where half of the mass lies in particles with a diameter less than the MMAD and half in particles with diameters greater than the MMAD

3.114**material's compatibility with skin**

characteristic of material resulting in insignificant irritation to the wearers' skin when in direct contact

3.115**maximum flow condition**

those factors appropriate to the design specified by the manufacturer which give rise to the highest flow rate

3.116**maximum use concentration****MUC**

maximum atmospheric concentration of a hazardous substance from which the RPD wearer can be expected to be protected when wearing an RPD, and is determined by the protection level of the respiratory protective device or class of respiratory protective devices and the occupational exposure limit of the hazardous substance

NOTE The maximum use concentration usually can be determined mathematically by multiplying the protection level specified for a respiratory protective device by the occupational exposure limit used for the hazardous substance.

3.117**mechanical shock**

dynamic force on the device during a fall from a given height as determined by a laboratory test

3.118**mechanical strength of visor**

ability of the device to withstand mechanical stress to the barrier in front of the eyes

3.119**medium pressure**

pressure between 100 hPa (mbar) above atmospheric pressure to 1 MPa (10 000 mbar) pressure gauge

3.120

metabolic simulator

programmable automatic **breathing machine** that can simulate the characteristics of both human breathing (variable tidal volume and respiratory rate, humidity, temperature) and metabolic functions (variable oxygen consumption and carbon dioxide production)

3.121

minimum flow condition

those factors appropriate to the design specified by the manufacturer which give rise to the lowest flow rate

3.122

minimum required protection factor

MRPF

level of protection required, derived from the **adequacy assessment** procedure

3.123

minute ventilation

total volume of air exchanged in the lungs during one minute, V_E in l/min **BTPS**

3.124

minute volume

quantity of air exhaled in one minute

3.125

mist

liquid droplets generated by condensation of vapour back to the liquid state or by breaking up a liquid into a dispersed state

3.126

mode of operation

primary means of supplying the wearer with breathable gas, i.e., particle filtering, gas and vapour filtering, or breathable gas supplying

3.127

monitor

component of an **RPD** to enable the wearer to continuously assess that the manufacturer's minimum design air flow rate or manufacturer's minimum design conditions are met

3.128

multi-functional RPD

RPD which are capable of operating within its mode of operation by using different operating methods; e.g. power assist filtering devices, power on/off, compressed **breathable gas** system with compressed breathable gas device

3.129

multi-type combination filter

combination filter which meets the requirements of more than one type of gas and particle filter

3.130

multi-type gas filter

gas filter which meets the requirements of more than one type of gas filter

3.131

negative pressure

pressure inside the respiratory interface, hose, etc. that is lower than that of the ambient atmosphere

NOTE

It is expressed in pascals (Pa).

3.132

nominal working duration

working time of a device, used for the classification determined in laboratory tests with a flow rate specified in an International Standard

3.133

occupational exposure limit

OEL

maximum concentration of airborne contaminants deemed to be acceptable, as defined by the authority having jurisdiction

3.134

operating pressure

pressure(s) developed within the **RPD** during service

3.135

oxygen compatibility

capability of **RPD** to allow direct contact with pressurised oxygen without risk of fire or explosion by being oil and grease free

3.136

oxygen consumption

amount of oxygen consumed by the human tissues for aerobic energy production, VO_2 in l/min **STPD**

3.137

oxygen deficient

condition based on an oxygen concentration or partial pressure below which a person may be adversely affected

NOTE

Each authority having jurisdiction may establish an alternative definition or specific limit.

3.138

oxygen-enriched air

air containing oxygen at higher concentrations than that of atmospheric air at sea level

3.139**particle**

generic term used to describe solid or liquid substances in the finely divided state and refers to particulate aerosols

EXAMPLE Dusts, mists, smoke, fumes, fibres, and fog as well as micro-organisms.

3.140**particle filter**

filter which is intended to remove airborne particles

3.141**particle filter efficiency**

degree to which a filter removes **aerosols** from the **ambient atmosphere**

3.142**particle filter respiratory protective device****particle filter RPD**

device consisting of a **respiratory interface** with a **particle filter** that removes finely divided **particles** from the air to be inhaled by the **wearer**

NOTE The filter medium may be replaceable or be an integral part of the construction.

3.143**peak inspiratory flow rate**

highest instantaneous flow rate during the inhalation phase of a breath cycle in l/s **BTPS**

NOTE Litres per second (l/s) is the preferred unit as the flow takes place only during a short fraction of the breath cycle.

3.144**permeation**

process by which a chemical moves through an **RPD** material on a molecular level (diffusion)

3.145**physical work capacity**

ability of a person to engage in muscular work

3.146**positive pressure**

pressure inside the **respiratory interface**, hose, etc. higher than that of the **ambient atmosphere**

3.147**practical performance**

evaluation of **RPD** during simulation of typical work or escape activities in a laboratory

3.148**protection level**

degree of respiratory protection allocated to an **RPD** for the purposes of selection and use that is expected to be provided to wearers when used within an effective **RPD** program as described in ISO 16975

3.149**pressure dew point**

temperature at which moisture begins to condense from a gas under pressure, as the gas is cooled at this pressure

3.150**psychological impact on wearer**

positive and negative influences on the wearer's state of mind by wearing the **RPD** or by its appearance and/or design

3.151**qualitative fit factor****QLFF**

minimum assured (C_o/C_i)

qualitative estimate of the minimum fit of a particular **tight-fitting respiratory protective device** to a specific individual when a **qualitative fit test** is passed, i.e., the test agent is not detected by the subject's senses

3.152**qualitative fit test****QLFT**

pass/fail test method that relies on the subject's sensory response to detect a challenge agent in order to assess the adequacy of **RPD** fit

3.153**quantitative fit factor****QNFF**

$QNFF = (C_o/C_i)$

measure of the fit of a particular **tight-fitting** respiratory interface to a specific individual

NOTE It represents only **respiratory interface** to face leakage. Leakage from other sources (e.g., air-purifying elements) must be essentially zero. The **QNFF** is measured with specialized instrumentation.

3.154**quantitative fit test****QNFT**

test method that uses an instrument to assess (quantify) the amount of face seal leakage into the **respiratory protective device** in order to assess the adequacy of its fit