

इथाइल एक्रिलेट — विशिष्टि
(पहला पुनरीक्षण)

Ethyl Acrylate — Specification
(*First Revision*)

ICS 71.080.70

© BIS 2024



भारतीय मानक ब्यूरो
BUREAU OF INDIAN STANDARDS
मानक भवन, 9 बहादुर शाह ज़फर मार्ग, नई दिल्ली - 110002
MANAK BHAVAN, 9 BAHADUR SHAH ZAFAR MARG
NEW DELHI - 110002
www.bis.gov.in www.standardsbis.in

FOREWORD

This Indian Standard (First Revision) was adopted by the Bureau of Indian Standards, after the draft finalized by the Organic Chemicals, Alcohols and Allied Products Sectional Committee had been approved by the Petroleum, Coal and Related Products Division Council.

Ethyl acrylate is produced by the esterification of acrylic acid or by acrylonitrile. It has versatile use in the manufacturing of leather auxiliaries, paper coating etc. It also finds its uses as a textile sizing agent and kerosene substitute water-based thickener.

Ethyl acrylate is a flammable liquid. Its vapours can form explosive mixtures with air and can cause irritation. It should be kept away from heat sources and chemicals that can cause its polymerization violently evolving considerable heat.

This standard was first published in 1999 by taking considerable assistance from the following standards:

- ASTM D 1364-1980 Test method for water in volatile solvents
- ASTM D 1613-1985 Test method for acidity in volatile solvents
- ASTM D 3125-1983 Test method for MEHQ in colourless monomeric acrylate esters and acrylic acids
- ASTM D 3362-1984 Test method for purity of acrylate esters by gas chromatography
- ASTM D 3548-1993 Specification for ethyl acrylate

In this (*first*) revision, requirement of colour has been modified to 10 Pt-Co from 20 Pt-Co and moisture content to 0.05 percent from 0.2 percent, as the limit present was high as compared to the material available in market. Also, the requirement of inhibitor has been modified to 10 ppm to 20 ppm, as inhibitor is a must to avoid polymerization and thus minimum amount of inhibitor is incorporated. Further, instrumental methods for determination of moisture content, colour and inhibitor content have been incorporated.

The Composition of the Committee responsible for formulation of this standard is given in [Annex E](#).

For the purpose of deciding whether a particular requirement of this standard is complied with, the final value, observed or calculated, expressing the result of a test or analysis, shall be rounded off in accordance with IS 2 : 2022 'Rules for rounding off numerical values (*second revision*)'. The number of significant places retained in the rounded off value should be the same as that of the specified value in this standard.

Indian Standard
ETHYL ACRYLATE — SPECIFICATION
(First Revision)

1 SCOPE

This standard prescribes the requirements, the methods of sampling and tests for ethyl acrylate.

2 REFERENCES

The standards given below contain provisions which through reference in this text, constitute provision of this standard. At the time of publication, the editions indicated were valid. All standards are subject to revision and parties to agreements based on this standard are encouraged to investigate the possibility of applying the most recent edition of these standards:

<i>IS No.</i>	<i>Title</i>
IS 1070 : 2023	Reagent grade water — Specification (<i>fourth revision</i>)
IS 1448	Methods of test for petroleum and its products:
(Part 178) : 2020/ ISO 6271 : 2015	Clear liquids — Estimation of colour by the platinum cobalt colour scale
(Part 182) : 2020/ ISO 12937 : 2000	Petroleum products — Determination of water — Coulometric Karl Fischer titration method
IS 2362 : 1993	Determination of water by Karl Fischer method — Test method (<i>second revision</i>)
IS 4905 : 2015/ ISO 24153 : 2009	Random sampling and randomization procedures (<i>first revision</i>)
IS 8768 : 2000	Method of measurement of colour in liquid chemical products platinum-cobalt scale (<i>second revision</i>)

3 REQUIREMENTS

3.1 The material shall also comply with the requirements given in [Table 1](#), when tested

according to the methods referred in col (4) and col (5) of [Table 1](#).

3.2 Quality of Reagents

Unless specified, otherwise, pure chemicals and distilled water (*see* 1070) shall be employed in tests.

NOTE — 'Pure chemicals' shall mean chemicals that do not contain impurities which affect the results of analysis.

4 PACKING AND MARKING**4.1 Packing**

The material shall be packed as agreed to between the supplier and the purchaser.

4.2 Marking

4.2.1 The material shall be legibly marked with the following information:

- a) Name of the material;
- b) Name of the manufacturer and its trade mark, if any;
- c) Month and year of manufacture;
- d) Batch number, in code or otherwise;
- e) Net mass of the material; and
- f) Any other statutory requirement.

4.2.2 BIS Certification Marking

The product(s) conforming to the requirements of this standard may be certified as per the conformity assessment schemes under the provisions of the *Bureau of Indian Standards Act, 2016* and the rules and regulations framed thereunder, and the products may be marked with the Standard Mark.

5 SAMPLING

The method of drawing representative samples shall be as prescribed in [Annex D](#).

To access Indian Standards click on the link below:

https://www.services.bis.gov.in/php/BIS_2.0/bisconnect/knowyourstandards/Indian_standards/isdetails/

Table 1 Requirements for Ethyl Acrylates(Clauses [3.1](#) and [D-5](#))

SI No.	Characteristic	Requirement	Method of Test, Ref to	
			Annex (4)	IS (5)
(1)	(2)	(3)		
i)	Assay, percent by mass, <i>Min</i>	99.5	A	—
ii)	Acidity (as acrylic acid), percent by mass, <i>Max</i>	00.01	B	—
iii)	Colour, Pt-Co scale, <i>Max</i>	10.00	—	IS 8768/IS 1448 (Part 178) ¹⁾
iv)	Water, percent by mass, <i>Max</i>	00.05	—	IS 2362/IS 1448 (Part 182) ¹⁾
v)	Inhibitors (as monomethyl ether hydroquinone), ppm, <i>Max</i>	10 to 20	C	—

¹⁾ In case of dispute, IS 1448 (Part 178) shall be the referee method for determination of colour and IS 1448 (Part 182) shall be the referee method for determination water.

ANNEX A

[Table 1, Sl No. (i)]

DETERMINATION OF ETHYL ACRYLATE CONTENT (ASSAY)

A-1 GENERAL

This test method covers the determination of the purity of ethyl acrylate by gas chromatography and, in addition, provides a means for measuring certain impurities such as alcohols and other esters. Water and acidity are measured by other methods and the results are used to normalize the chromatographic values.

A-2 APPARATUS

A-2.1 Gas Chromatograph

A-2.1.1 Any suitable gas liquid chromatographic instrument with a flame ionization detector may be used with following accessories and typical operating conditions:

Column length	6 m (20 ft)
Internal diameter	6.4 mm (1/4 inch)
Syringe	2 µl
Sample size	1 µl to 2 µl
Recorder	A recording potentiometer with a full scale deflection of 1 mV, full scale response time of 2 s or less
Carrier gas	Helium of purity 99.95 mol percent or nitrogen gas
Column material	Copper/stainless steel/aluminum
Liquid phase	Polyalkylene glycol (PAG) or equivalent phase capable of achieving desired separation of various components of interest
Solid support	Chromosorb P-acid washed 45 to 60 mesh size
Solvent	Methylene chloride, reagent grade

A-2.2.2 Instrument Conditions

	Temperature, °C
Column, isothermal	100
Injection port	200
Detector block	260
Carrier gas	Helium
Flow rate, ml/min	6.0
Typical retention time, min	17.7

NOTE — The above gas chromatographic conditions are suggestive. However, any GC having different columns (packed/capillary having different length/diameter/film thickness) and different carrier gas (He, H₂ or N₂), with different calibration technique (internal standard, external standard, area normalization) may be used provided standardization/calibrations are done after setting up chromatographic conditions for required resolution.

A-3 STANDARDS FOR CALIBRATION AND IDENTIFICATION

Standard samples of all components present are needed for identification by retention time and for calibration of quantitative measurement.

A-4 PROCEDURE

Introduce a representative specimen into the chromatograph using sufficient specimen to ensure a minimum 10 percent recorder deflection for a 0.1 percent concentration of purity at the most sensitive setting of instrument. Using the same conditions as for component identification and standardization, record the peaks of all components at attenuation setting that provide maximum peak heights.

A-5 CALCULATION

A-5.1 Measure the area of all peaks and multiply each area by the appropriate detector response factors to correct the difference in response to the components. Calculate the weight percent composition by dividing the individual corrected component areas by the total corrected area. Make corrections to account for water and acidity present.

A-5.2 Calculate the percent by mass of each component as follows:

$$\text{Component content, percent by mass} = \frac{A \times (100 - C)}{B}$$

where

A = corrected peak response;

B = sum of corrected peak responses; and

C = sum of water and acidity.

A-6 PRECISION AND BIAS

A-6.1 Repeatability

Two results, should be considered suspect if they differ by more than 0.06 percent.

A-6.2 Reproducibility

Two results should be considered suspect if they differ by more than 0.27 percent.

ANNEX B

[Table 1, Sl No. (ii)]

DETERMINATION OF ACIDITY

B-1 GENERAL

This test method covers the determination of total acidity as acrylic acid, in concentrations below 0.05 percent in ethyl acrylate. The specimen is mixed with either an equal volume of water or an equal volume of alcohol and titrated with aqueous sodium hydroxide solution to the phenolphthalein end point.

B-2 APPARATUS

B-2.1 Burette — 10 ml, graduated in 0.05 ml

B-2.2 Erlenmeyer Flask — 250 ml

B-3 REAGENTS

B-3.1 Alcohols — refined ethyl or iso-propyl

B-3.2 Phenolphthalein Indicator Solution — 10 g/l

Dissolve 1 g of phenolphthalein in ethyl or isopropyl alcohol and dilute to 100 ml with the alcohol.

NOTE — Bromophenol blue indicator may also be used.

B-4 PROCEDURE

Measure into a 250 ml conical flask 50 ml of water, if the sample is water soluble, or 50 ml of alcohol, if

the sample is not completely water soluble. Add 0.5 ml of phenolphthalein indicator solution. Titrate with water/alcohol with 0.05 N sodium hydroxide (NaOH) solution to the first pink colour. Pipette 50 ml of the sample into the flask. Titrate with the 0.05 N sodium hydroxide (NaOH) solution to the same first pink colour originally obtained.

B-5 CALCULATION

Acidity as acrylic acid, percent by mass =

$$\frac{V \times N \times 0.144}{D}$$

where

V = sodium hydroxide (NaOH) solution required for titration of the sample, in ml;

N = normality of sodium hydroxide (NaOH); and

D = specific gravity of the specimen at the test temperature.

B-6 PRECISION AND BIAS

Repeatability 0.000 8 percent

Reproducibility 0.001 4 percent

NOTE — Auto Titrator may also be used for determination of acidity.

ANNEX C

[Table 1, Sl No. (v)]

DETERMINATION OF INHIBITORS

C-1 GENERAL

Two methods namely, Method A for determination of inhibitor content by spectrophotometer and Method B for determination of inhibitor content by gas chromatography have been prescribed. In case of dispute, Method B shall be the referee method.

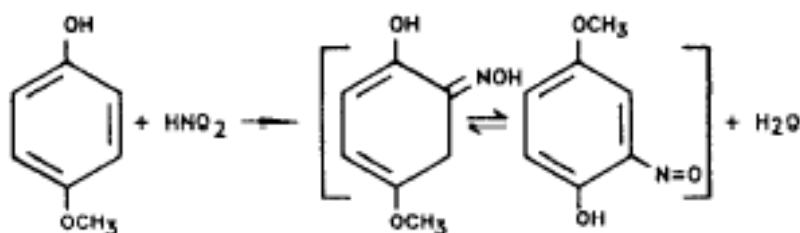
C-2 METHOD A

C-2.1 General

This test method covers the determination of

monomethyl ether of hydroquinone (MEHQ) in ethyl acrylate. It is applicable to the determination of MEHQ in the concentration range from 0 ppm to 1 200 ppm. MEHQ reacts with nitrous acid (sodium nitrite in acidic media) to form the nitroso derivative which equilibrates between two structures as given below.

The yellow colour of the nitroso compound is measured spectrophotometrically at a wavelength of 420 nm.



C-2.2 Apparatus

C-2.2.1 Spectrophotometer — with borosilicate glass cells for determining absorbance at 420 nm

C-2.2.2 Volumetric Flasks — 50 ml and 100 ml capacity

C-2.2.3 Measuring Pipette — 5 ml and 10 ml capacity

C-2.3 Reagents

C-2.3.1 Acetic Acid — glacial

C-2.3.2 Methyl Ether of Hydroquinone (MEHQ) (4-Methoxy Phenol)

C-2.3.3 Sodium Nitrite Solution

Dissolve 2 g of sodium nitrite in water and dilute to 100 ml.

C-2.4 Calibration

C-2.4.1 Weigh 0.10 g of MEHQ to the nearest 0.1 mg into a 100 ml volumetric flask containing approximately 50 ml of glacial acetic acid. Mix well until solution is complete then dilute to the mark with glacial acetic acid. Prepare a series of standards by pipetting 1 ml, 2 ml, 4 ml, 6 ml and 10 ml portions of the MEHQ solution into respective 50 ml volumetric flasks. Dilute each flask to the mark with glacial acetic acid and mix well. A 10 ml aliquot of each of these standards contains 200 µg, 400 µg, 800 µg, 1 200 µg and 2 000 µg of MEHQ respectively.

C-2.4.2 Determine the absorbance each of these standards by pipetting 10 ml aliquots into 50 ml volumetric flasks containing 20 ml of glacial acetic acid. To each flask add 1 ml of 2 percent sodium nitrite (NaNO_2), solution and dilute to the mark with glacial acetic acid. Mix well and allow to stand for 10 min. With cells appropriate to the instrument, determine the absorbance at 420 nm using acetic acid as the blank. Construct a calibration curve on

rectangular coordinate graph paper by plotting the absorbances of the standards at 420 nm against the micrograms of MEHQ.

C-2.5 Procedure

C-2.5.1 Perform analysis in duplicate and carry a blank through the analysis using 49 ml of glacial acetic acid in place of the specimen solution. Weigh the appropriate amount of specimen into a 50 ml volumetric flask containing 20 ml of glacial acetic acid.

C-2.5.2 Add 1 ml of 2 percent sodium nitrite (NaNO_2) solution to the specimen and dilute to the mark with glacial acetic acid. Mix well and allow to stand for 10 min.

C-2.5.3 Using the procedure followed for the calibration, determine the absorbance of the solution at 420 nm with the blank solution in the reference position. From the calibration curve, determine the micrograms of MEHQ corresponding to the absorbance obtained.

C-2.6 Calculation

Calculate the concentration of MEHQ in ppm as follows:

$$\text{MEHQ, ppm} = \frac{M}{S}$$

where

M = amount of MEHQ from calibration curve, in µg; and

S = amount of specimen used in the test, in g.

C-2.7 Precision

C-2.7.1 Repeatability

Two results, each the mean of duplicates, obtained by the same operator on different days should be considered suspect if they differ by more than 4.0 percent relative.

C-2.7.2 Reproducibility

Two results, each the mean of duplicates, obtained by operators in different laboratories should be considered suspect if they differ by more than 15 percent relative.

C-3 METHOD B**C-3.1 Apparatus****C-3.1.1 Gas Chromatograph**

C-3.1.1.1 Any gas chromatograph equipped with a flame ionization detector (FID), a split/splitless injector and a suitable electronic integrator/software, capable of operating at the conditions listed below, may be used:

Column	:	Stationary phase of (5 percent phenyl)-methylpolysiloxane with length 30 m; internal diameter 0.32 mm and film thickness 1.0 µm or equivalent
Sample size	:	1 µl
Flow		
Carrier gas	:	Helium at constant flow of 1 ml/min or nitrogen gas
Air flow rate	:	450 ml/min
Hydrogen flow rate	:	40 ml/min
Make-up gas	:	Nitrogen or helium at constant flow rate of 30 ml/min
Split ratio	:	100 : 1
Run time	:	39 min

C-3.1.1.2 Temperature Programme of Oven, Detector and Injector:

Sl No.	Injector Temperature, °C	Detector Temperature, °C	Oven		
			Temperature, °C	Hold Time, min	Ramp Rate, °C/min
(1)	(2)	(3)	(4)	(5)	(6)
i)	250	300	50	5	—
			110	—	4

NOTE — The above gas chromatographic conditions are suggestive. However, any GC having different columns (packed/capillary having different length/diameter/film thickness) and different carrier gas (He, H₂ or N₂), with different calibration technique (internal standard, external standard, area normalization) may be used provided standardization/calibrations are done after setting up chromatographic conditions for required resolution.

C-3.1.2 Analytical Balance**C-3.1.3 Vial with screw cap — 30 ml****C-3.1.4 Autosampler — 2 ml**

NOTE — Syringe may also be used for injecting the sample.

C-3.2 Reagents

C-3.2.1 Ethyl Acrylate — purity of 99.5 percent, minimum

C-3.2.2 Ethyl Acetate — purity of 99.5 percent, minimum

C-3.2.3 Monomethyl ether of hydroquinone (MEHQ, 4-methoxyphenol) — purity of 99 percent, minimum

C-3.3 Procedure

Inject 1 µl aliquot of the sample manually by syringe or by automatic sampler into the column and obtain the chromatograph. Determine the mass concentration of all components by area normalization method.

C-3.4 Calculation

MEHQ content, percent by mass

$$= \frac{R_{MEHQ} \times A_{MEHQ}}{\sum(R_i \times A_i)}$$

where

R_{MEHQ} = relative response factor of MEHQ;

R_i = relative response factor of each component;

A_{MEHQ} = peak area of MEHQ;

A_i = peak area of each component; and

$\sum(R_i \times A_i)$ = sum of product of each peak area times its relative response factor.

C-3.5 Report

Report inhibitors (as monomethyl ether hydroquinone) content in ppm.

ANNEX D

(Clause 5)

SAMPLING OF ETHYL ACRYLATE

D-1 GENERAL

D-1.1 Samples shall be taken at a place protected from damp air, dust and soot.

D-1.2 Sampling instrument shall be clean and dry.

D-1.3 Precautions shall be taken to protect the samples, the material being sampled, the sampling instrument and the containers for samples from adventitious contamination.

D-1.4 To draw a representative sample, the contents of each container, selected for sampling, shall be mixed, as thoroughly as possible, by suitable means.

D-1.5 The samples shall be placed in clean, dry and air-tight glass or other suitable containers on which the material has no action.

D-1.6 The sample containers shall be of such a size that they are almost completely filled by the sample.

D-1.7 Each sample container shall be sealed air-tight after filling and marked with full details of sampling, the date of sampling, year of manufacture and other important particulars of the consignment.

D-1.8 Samples shall be stored in a cool and dry place.

D-2 SCALE OF SAMPLING**D-2.1 Lot**

All the containers in a single consignment of the material drawn from a single batch of manufacture shall constitute a lot. If a consignment is declared or known to consist of containers pertaining to different batches of manufacture, the containers belonging to the same batch of manufacture shall be grouped together and each such group shall constitute a separate lot.

D-2.2 For ascertaining the conformity of the lot to the requirements of this specification, tests shall be carried out for each lot separately.

D-2.3 The number of containers (n) to be selected for drawing the samples shall depend upon the size of the lot (N) and shall be in accordance with [Table 2](#).

D-2.4 These containers shall be selected at random from the lot and to ensure the randomness of selection, procedure given in IS 4905 may be followed.

D-3 TEST SAMPLE AND REFEREE SAMPLE

D-3.1 From each of the container selected as in [D-2.3](#), draw with an appropriate sampling instrument small portions of the material from different parts of the container. The total quantity so drawn from each of the containers shall be approximately equal to thrice the quantity required for testing purposes.

D-3.2 Mix thoroughly all the portions of the material drawn from the same container to give a representative sample for the container.

D-3.3 From the samples (*see* [D-3.2](#)) representing different containers selected in [D-2.3](#), a small but equal quantity of material shall be taken and thoroughly mixed to form a composite sample, sufficient to carry out testing for the characteristics specified. The composite sample obtained shall be divided into three equal parts, one for the purchaser, another for the supplier and the third for the referee.

D-3.4 The referee samples consisting of a composite sample shall bear the seals of both the purchaser and the supplier and shall be kept at a place agreed to between the two. This shall be used in case of any dispute between the two.

D-4 TESTS

Tests for all the characteristics shall be conducted on the composite sample.

D-5 CRITERIA FOR CONFORMITY

For declaring the conformity of a lot to the requirements of all other characteristics (*see* [D-4](#)) tested on the composite sample, the test results for the characteristics shall satisfy the relevant requirements given in [Table 1](#).

Table 2 Number of Containers to be Selected for Sampling*(Clause [D-2.3](#))*

Sl No.	Lot Size	No. of Containers to be Selected
	<i>(N)</i>	<i>(n)</i>
(1)	(2)	(3)
i)	Up to 25	2
ii)	26 to 50	3
iii)	51 to 100	5
iv)	101 to 300	7
v)	301 and above	10

NOTE — When the size of the lot is three or less, all the containers shall be sampled.

ANNEX E

(Foreword)

COMMITTEE COMPOSITION

Organic Chemicals, Alcohols and Allied Products Sectional Committee, PCD 09

<i>Organization</i>	<i>Representative(s)</i>
National Chemical Laboratory (NCL), Pune	DR C. V. RODE (<i>Chairperson</i>)
All India Distillers Association (AIDA), New Delhi	SHRI SUKHRAJ SONI SHRI A. K. SINGHAL (<i>Alternate I</i>) SHRI RAJESH DHINGRA (<i>Alternate II</i>)
BASF India Limited, Mumbai	SHRI DATTATRAY ANNASO GURAV SHRI HEMAL (<i>Alternate</i>)
Chemical and Petrochemicals Manufacturers Association (CPMA), New Delhi	SHRI UDAY CHAND
CSIR - Central Drug Research Institute (CDRI), Lucknow	DR SANJEEV KANOJIYA
Deepak Fertilizers and Petrochemicals Corporation Limited, Navi Mumbai	SHRI SURESH AMLE DR L. B. YADAWA (<i>Alternate</i>)
Deepak Phenolics Limited, Vadodara	SHRI DHARMESH SIDDHAPURIA SHRI MEHUL KUMAR PATEL (<i>Alternate</i>)
Dow Chemical International Private Limited, Mumbai	SHRI V. MOHANDOSS SHRI GOVIND GUPTA (<i>Alternate</i>)
Godavari Biorefineries, Mumbai	SHRI SHANUL LAXMANRAO PAGAR SHRI APPASAHEB J. WANI (<i>Alternate</i>)
Gujarat Narmada Valley Fertilizers Company Limited, Ahmedabad	SHRI R. M. PATEL SHRI C. S. PATEL (<i>Alternate</i>)
Hindustan Organic Chemicals Limited (HOCL), Mumbai	DR B. RAJEEV
India Glycols Limited, Uttarakhand	DR R. K. SHARMA SHRI ALOK SINGHAL (<i>Alternate</i>)
Indian Chemical Council (ICC), Mumbai	SHRI J. I. SEVAK SHRI DHRUMIL SONI (<i>Alternate</i>)
Indian Oil Corporation Limited, Panipat	SHRI Y. S. JHALA
Jubilant Agri and Consumer Products Limited, Gurugram	SHRI KANAK BARAN DASS
Laxmi Organic Industries, Mumbai	SHRI KRISHNA A. RAO SHRI KAMLESH FULCHAND SHINDE (<i>Alternate</i>)

<i>Organization</i>	<i>Representative(s)</i>
Ministry of Chemicals and Fertilizers, New Delhi	SHRI O. P. SHARMA SHRI VARUN SINGH POONIA (<i>Alternate</i>)
National Chemical Laboratory (NCL), Pune	DR RAVINDAR KONTHAM DR UDAYA KIRAN MARELLI (<i>Alternate</i>)
Reliance India Limited (RIL), Mumbai	SHRI K. K. SREERAMACHANDRAN SHRI VASANT WARKE (<i>Alternate</i>)
United Phosphorus Limited (UPL), Mumbai	SHRI M. D. VACHHANI
In Personal Capacity (<i>37 Nandanvan Society, Near GNFC Township, Narmadanagar, Bharuch – 392015</i>)	DR MAYUR J. KAPADIA
BIS Directorate General	SHRI CHINMAY DWIVEDI, SCIENTIST ‘E’/DIRECTOR AND HEAD (PETROLEUM, COAL AND RELATED PRODUCTS) [REPRESENTING DIRECTOR GENERAL (<i>Ex-officio</i>)]

Member Secretary
MS ADITI CHOUDHARY
SCIENTIST ‘C’/DEPUTY DIRECTOR
(PETROLEUM, COAL AND RELATED PRODUCTS), BIS

Bureau of Indian Standards

BIS is a statutory institution established under the *Bureau of Indian Standards Act, 2016* to promote harmonious development of the activities of standardization, marking and quality certification of goods and attending to connected matters in the country.

Copyright

BIS has the copyright of all its publications. No part of these publications may be reproduced in any form without the prior permission in writing of BIS. This does not preclude the free use, in the course of implementing the standard, of necessary details, such as symbols and sizes, type or grade designations. Enquiries relating to copyright be addressed to the Head (Publication & Sales), BIS.

Review of Indian Standards

Amendments are issued to standards as the need arises on the basis of comments. Standards are also reviewed periodically; a standard along with amendments is reaffirmed when such review indicates that no changes are needed; if the review indicates that changes are needed, it is taken up for revision. Users of Indian Standards should ascertain that they are in possession of the latest amendments or edition by referring to the website-www.bis.gov.in or www.standardsbis.in.

This Indian Standard has been developed from Doc No.: PCD 09 (24540).

Amendments Issued Since Publication

Amend No.	Date of Issue	Text Affected

BUREAU OF INDIAN STANDARDS

Headquarters:

Manak Bhavan, 9 Bahadur Shah Zafar Marg, New Delhi 110002

Telephones: 2323 0131, 2323 3375, 2323 9402

Website: www.bis.gov.in

Regional Offices:

Central : 601/A, Konnectus Tower -1, 6th Floor,
DMRC Building, Bhavbhuti Marg, New
Delhi 110002

Telephones

{ 2323 7617

Eastern : 8th Floor, Plot No 7/7 & 7/8, CP Block, Sector V,
Salt Lake, Kolkata, West Bengal 700091

{ 2367 0012
{ 2320 9474

Northern : Plot No. 4-A, Sector 27-B, Madhya Marg,
Chandigarh 160019

{ 265 9930

Southern : C.I.T. Campus, IV Cross Road, Taramani, Chennai 600113

{ 2254 1442
{ 2254 1216

Western : 5th Floor/MTNL CETTM, Technology Street, Hiranandani Gardens,
Powai, Mumbai 400076

{ 2570 0030
{ 2570 2715

Branches : AHMEDABAD, BENGALURU, BHOPAL, BHUBANESHWAR, CHANDIGARH, CHENNAI, COIMBATORE, DEHRADUN, DELHI, FARIDABAD, GHAZIABAD, GUWAHATI, HARYANA (CHANDIGARH), HUBLI, HYDERABAD, JAIPUR, JAMMU, JAMSHEDPUR, KOCHI, KOLKATA, LUCKNOW, MADURAI, MUMBAI, NAGPUR, NOIDA, PARWANOO, PATNA, PUNE, RAIPUR, RAJKOT, SURAT, VIJAYAWADA.