

ETHYL ALCOHOL

Prepared at the 46th JECFA (1996), published in FNP 52 Add 4 (1996) superseding specifications prepared at the 29th JECFA (1985), published in FNP 34 (1986. An ADI limited by GMP was established at 14th JECFA (1970))

SYNONYMS

Alcohol, ethanol

DEFINITION

Contains usually about 95% v/v (i.e. with about 5% v/v water) C₂H₆O; other grades with a different water content may be used depending on the technological requirements; these specifications apply only to undenaturated ethyl alcohol.

Chemical names

Ethanol

C.A.S. number

64-17-5

Chemical formula

C₂H₆O

Structural formula



Formula weight

46.07

Assay

Not less than 94.9% v/v

DESCRIPTION

Clear colourless mobile liquid with a mild characteristic odour; flammable

FUNCTIONAL USES

Extraction solvent, carrier solvent, flavouring agent (see "Flavouring agents" monograph JECFA no. 41)

CHARACTERISTICS

IDENTIFICATION

Solubility (Vol. 4)

Soluble in water

Refractive index (Vol. 4)

n (20, D): 1.364

Boiling point (Vol. 4)

78°

Infrared absorption

The infrared spectrum of the sample corresponds with the reference infrared spectrum below

PURITY

Solubility

Transfer 50 ml of the sample to a 100 ml glass-stoppered graduated cylinder, dilute to 100 ml with water, and mix. Place the cylinder in a water bath maintained at 10°, and allow to stand for 30 min. No haze or turbidity develops.

Non-volatile residue
(Vol. 4)

Not more than 2 mg/100 ml

<u>Acidity</u>	Not more than 0.005% w/v as acetic acid Transfer 10 ml of the sample to a glass-stoppered flask containing 25 ml of water, add 0.5 ml of phenolphthalein TS. Add 0.02N sodium hydroxide until the first appearance of a pink colour that persists after shaking for 30 sec. Add 25 ml (about 20 g) of the sample, mix, and titrate with 0.02N sodium hydroxide until the pink colour is restored. Not more than 1.0 ml is required.
<u>Alkalinity</u>	Not more than 0.003% as ammonia Add 2 drops of methyl red TS to 25 ml of water. Add 0.02N sulfuric acid until a red colour just appears, then add 25 ml (about 20 g) of the sample, and mix. Not more than 2.0 ml of 0.02N sulfuric acid is required to restore the red colour.
<u>Fusel oil</u>	Mix 10 ml of the sample with 1 ml of glycerol and 1 ml of water, and allow to evaporate from a piece of clean, odourless absorbent paper. No foreign odour is perceptible when the last traces of ethanol leave the paper.
<u>Ketones and other alcohols</u>	Total: Not more than 0.5% Methanol: Not more than 0.02% Other: Not more than 0.1% See description under TESTS
<u>Readily carbonizable substances</u>	Transfer 10 ml of sulfuric acid into a small Erlenmeyer flask, cool to 10°, and add 10 ml of the sample, dropwise, with constant agitation. The mixture is colourless or has no more colour than either the acid or sample before mixing.
<u>Readily oxidizable substances</u>	Transfer 20 ml of the sample, previously cooled to 15° to a glass-stoppered cylinder, add 0.1 ml of 0.1N potassium permanganate, mix and allow to stand for 5 min. The pink colour is not entirely discharged.
<u>Lead</u> (Vol. 4)	Not more than 0.5 mg/kg Determine using an atomic absorption technique appropriate to the specified level. The selection of sample size and method of sample preparation may be based on the principles of the method described in Volume 4, "Instrumental Methods."

TESTS

PURITY TESTS

<u>Ketones and other alcohols</u>	Proceed as directed under <i>Gas Liquid Chromatography</i> (Volume 4) using the following conditions: Column - length: 1.80 m - diameter: 6.4 mm - material: steel - packing: P.E.G. 400 10% - support: Chromosorb W 60/80 Carrier gas: Helium
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Flow rate: 45 ml/min

Detector type: FID

Temperatures

- injection port: 150°

- column: 90°

- detector: 150°

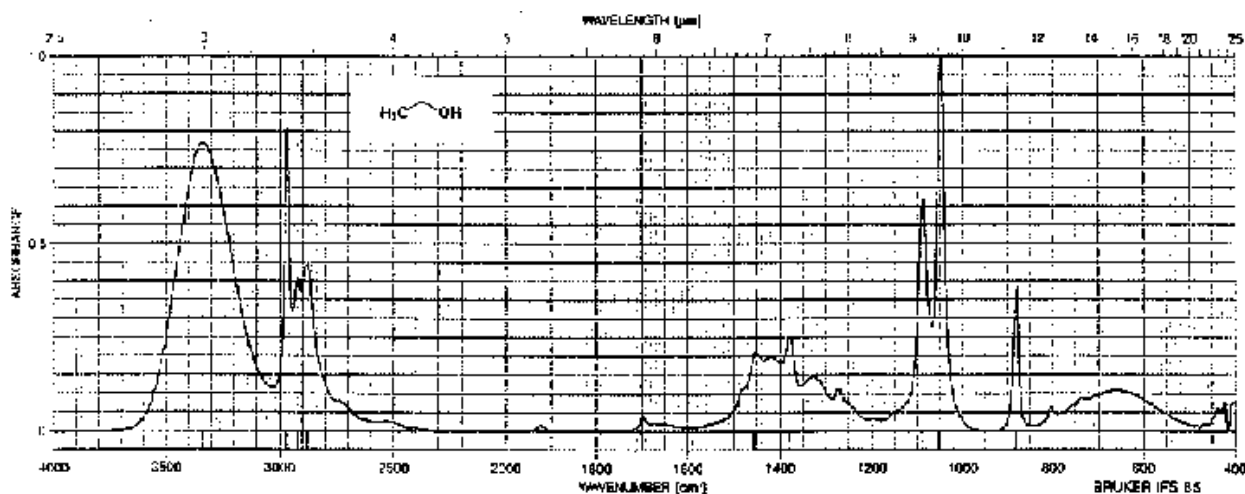
Inject 5 µl of the sample. The sum of areas of the peaks other than that of ethyl alcohol is not more than 0.5% of the sum of total peaks. The area of peak of methanol is not more than 0.02% and the area of any other individual peak of impurities is not more than 0.1% of the total.

METHOD OF ASSAY

The specific gravity of the sample is not higher than 0.8096 at 25°/25° (equivalent to 0.8161 at 15.56°/15.56°).

Infrared spectrum

Ethyl alcohol



Infrared spectrum from Merck FT-IR Atlas through courtesy of Dr. K.G.R. Pachler, Mr. F. Mutlok and Dr. H-U. Grenlich, c/o Merck, Darmstadt, and VCH Verlagsgesellschaft GmbH, Weinheim, Germany.