PHARMACOPOEIAL DISCUSSION GROUP

CORRECTION CODE: E48

NAME: ETHYL PARAHYDROXYBENZOATE

(Correction 2 to revision 1 of the sign-off document signed 10 June 2009)

Item to be corrected:

- Addition of CAS numbers: [120-47-8]
- Appearance of solution/color: addition of comparison with alcohol

Attribute	EP	JP	USP
Definition	+	+	+
Identification A	+	+	+
(melting point)*			
Identification B	+	+	+
(IR)			
Appearance of	+	+	+
solution/color			
Acidity	+	+	+
Related	+	+	+
substances**		5,	
Sulphated ash	+	+	+
Assay	+	+	+

^{*} Melting point: listed in JP as a test and not as part of identification

Legend

- + will adopt and implement
- will not stipulate

Non-harmonised attributes

Characters, Storage

Local requirements

Ph. Eur.	JP	USP
Second identification	Related substances: test for	none
(melting point, TLC)	required detectability, system	
	repeatability	
	Heavy metals (20 ppm)	
	Assay: column temperature	

Reagents and reference materials

Each pharmacopoeia will adapt the text to take account of local reference materials and reagent specifications.



^{**} Related substances: JP uses the term "relative response factor" instead of "correction factor"

European Pharmacopoeia

Signature <

Name

Date

VIELCE COME 25-200-209

Japanese Pharmacopoeia

Name

Date

Signature A Structure for Y. Yoshida

Havahiro ducla

16 Dec/2020

United States Pharmacopeia

Signature

Name

Date

JLT. M

KEVIN MOORE

19-NOV-2020

E48 - ETHYL PARAHYDROXYBENZOATE

C9H10O3

Mr 166.2

[120-47-8]

DEFINITION

Ethyl 4-hydroxybenzoate.

Content: 98.0 per cent to 102.0 per cent.

IDENTIFICATION

A. Melting point: 115 °C to 118 °C.

B. Infrared absorption spectrophotometry

Record the infrared absorption spectrum of ethyl parahydroxybenzoate and compare with the Reference Spectrum or the spectrum obtained with the Reference Standard: the transmission minima correspond in position and relative size.

TESTS

Solution S. Dissolve 1.0 g in *alcohol* and dilute to 10 ml with the same solvent.

Appearance of solution. Solution S is clear and not more intensely coloured than *alcohol* or the reference solution

Primary solutions:

- Ferric chloride primary solution: a 45.0 g/l solution of ferric chloride (FeCl₃, 6H₂O).
- Cobalt chloride primary solution: a 59.5 g/l solution of cobalt chloride (CoCl₂, 6H₂O).
- Copper sulphate primary solution: a 62.4 g/l solution of copper sulphate (CuSO₄, 5H₂O).

Reference solution:

To 5.0 ml of cobalt chloride primary solution, 12.0 ml of ferric chloride primary solution and 2.0 ml of copper sulphate primary solution, add hydrochloric acid (10 g/l HCl) to make 1000.0 ml.

Acidity. To 2 ml of solution S add 3 ml of alcohol, 5 ml of carbon dioxide-free water and 0.1 ml of bromocresol green solution. Not more than 0.1 ml of 0.1 M sodium hydroxide is required to change the colour of the indicator to blue.

Related substances. Liquid chromatography.

Test solution. Dissolve 50.0 mg of the sample to be examined in 2.5 ml of *methanol* and dilute to 50.0 ml with the mobile phase. Dilute 10.0 ml of this solution to 100.0 ml with the mobile phase.

W/Cm

Reference solution (a). Dissolve 5 mg each of methyl parahydroxybenzoate R, 4-hydroxybenzoic acid R and the substance to be examined in the mobile phase and dilute to 100.0 ml with the same solvent. Dilute 1 ml of this solution to 10.0 ml with the mobile phase.

Reference solution (b). Dissolve 50.0 mg of ethyl parahydroxybenzoate CRS in 2.5 ml of methanol and dilute to 50.0 ml with the mobile phase. Dilute 10.0 ml of this solution to 100.0 ml with the mobile phase.

Reference solution (c). Dilute 1.0 ml of the test solution to 20.0 ml with the mobile phase. Dilute 1.0 ml of this solution to 10.0 ml with the mobile phase.

Column:

— size: l = 0.15 m, $\emptyset = 4.6$ mm;

— stationary phase: octadecylsilyl silica gel for chromatography (5 μm).

Mobile phase: 6.8 g/l solution of potassium dihydrogen phosphate, methanol (35:65 V/V).

Flow rate: 1.3 ml/min.

Detection: 272 nm.

Injection: 10 µl of the test solution and reference solutions (a) and (c).

Run time: 4 times the retention time of ethyl parahydroxybenzoate.

Relative retention with reference to ethyl parahydroxybenzoate (retention time = about 3.0 min): 4-hydroxybenzoic acid = about 0.5; methyl parahydroxybenzoate = about 0.8.

System suitability:

- resolution: minimum of 2.0 between the peaks due to methyl parahydroxybenzoate and to ethyl parahydroxybenzoate in the chromatogram obtained with reference solution (a).

Limits:

- correction factor: for the calculation of content, multiply the peak area of 4-hydroxybenzoic acid by 1.4;
- 4-hydroxybenzoic acid: not more than the area of the principal peak in the chromatogram obtained with reference solution (c) (0.5 per cent);
- unspecified impurities: for each impurity, not more than the area of the principal peak in the chromatogram obtained with reference solution (c) (0.5 per cent);
- total: not more than twice the area of the principal peak in the chromatogram obtained with reference solution (c) (1.0 per cent);
- disregard limit: 0.2 times the area of the principal peak in the chromatogram obtained with reference solution (c) (0.1 per cent).

Sulphated ash: maximum 0.1 per cent, determined on 1.0 g.

ASSAY

Liquid chromatography as described in the test for related substances with the following modification.

Injection: test solution and reference solution (b).

System suitability:

- repeatability: maximum relative standard deviation of 0.85 per cent after 6 injections of the reference solution (b).

CU Km

Calculate the percentage content of $C_9H_{10}O_3$ in the sample to be examined from the peak areas in the chromatograms obtained with test solution and reference solution (b) and the declared content of *ethyl parahydroxybenzoate CRS*.

REAGENTS

Bromocresol green solution.

Dissolve 50 mg of *bromocresol green* in 0.72 ml of 0.1 M sodium hydroxide and 20 ml of alcohol and dilute to 100 ml with water.

Test for sensitivity. To 0.2 ml of the bromocresol green solution add 100 ml of carbon dioxide-free water. The solution is blue. Not more than 0.2 ml of 0.02 M hydrochloric acid is required to change the colour to yellow. Colour change: pH 3.6 (yellow) to pH 5.2 (blue).

