



Edition: BP 2025 (Ph. Eur. 11.6 update)

Emulsifying Cetostearyl Alcohol (Type B)



[General Notices](#)

(Ph. Eur. monograph 0802)

Action and use

Excipient.

Ph Eur

DEFINITION

Mixture of cetostearyl alcohol and sodium laurilsulfate. A suitable buffer may be added.

Content

- [cetostearyl alcohol](#): minimum 80.0 per cent (anhydrous substance);
- [sodium laurilsulfate](#): minimum 7.0 per cent (anhydrous substance).

CHARACTERS

Appearance

White or pale yellow, waxy mass, plates, flakes or granules.

Solubility

Soluble in hot water giving an opalescent solution, practically insoluble in cold water, slightly soluble in ethanol (96 per cent).

IDENTIFICATION

First identification: B, C, D.

Second identification: A, C, D.

A. Thin-layer chromatography ([2.2.27](#)).

Test solution (a) Dissolve 0.1 g of the substance to be examined in 10 mL of [trimethylpentane R](#), heating on a water-bath. Shake with 2 mL of [ethanol \(70 per cent V/V\) R](#) and allow to separate. Use the lower layer as test solution (b). Dilute 1 mL of the upper layer to 8 mL with [trimethylpentane R](#).

Test solution (b) Use the lower layer obtained in the preparation of test solution (a).

Reference solution (a) Dissolve 24 mg of [cetyl alcohol CRS](#) and 16 mg of [stearyl alcohol CRS](#) in 10 mL of [trimethylpentane R](#).

Reference solution (b) Dissolve 20 mg of [sodium laurilsulfate CRS](#) in 10 mL of [ethanol \(70 per cent V/V\) R](#), heating on a water-bath.

Plate [TLC octadecylsilyl silica gel F₂₅₄ plate R](#).

Mobile phase [water R](#), [acetone R](#), [methanol R](#) (20:40:40 V/V/V).

Application 10 µL.

Development Over 2/3 of the plate.

Drying In air.

Detection Spray with a 50 g/L solution of [phosphomolybdic acid R](#) in [ethanol \(96 per cent\) R](#); heat at 120 °C until spots appear (about 5 min).

Results:

— the 2 principal spots in the chromatogram obtained with test solution (a) are similar in position and colour to the principal spots in the chromatogram obtained with reference solution (a);

— 1 of the spots in the chromatogram obtained with test solution (b) is similar in position and colour to the principal spot in the chromatogram obtained with reference solution (b).

B. Examine the chromatograms obtained in the assay of cetostearyl alcohol.

Results The 2 principal peaks in the chromatogram obtained with the test solution are similar in retention time to the 2 principal peaks in the chromatograms obtained with reference solutions (a) and (b).

C. It gives a yellow colour to a non-luminous flame.

D. To 0.3 g add 20 mL of [anhydrous ethanol R](#) and heat to boiling on a water-bath with shaking. Filter the mixture immediately, evaporate to dryness and take up the residue in 7 mL of [water R](#). To 1 mL of the solution add 0.1 mL of a 1 g/L solution of [methylene blue R](#), 2 mL of [dilute sulfuric acid R](#) and 2 mL of [methylene chloride R](#) and shake. A blue colour develops in the lower layer.

TESTS

[Acid value \(2.5.1\)](#)

Maximum 2.0.

[Iodine value \(2.5.4, Method A\)](#)

Maximum 3.0.

Dissolve 2.00 g in 25 mL of [methylene chloride R](#).

[Saponification value \(2.5.6\)](#)

Maximum 2.0.

[Water \(2.5.12\)](#)

Maximum 3.0 per cent, determined on 2.50 g.

ASSAY

Cetostearyl alcohol

Gas chromatography (2.2.28).

Internal standard solution Dissolve 0.200 g of [1-nonadecanol CRS](#) in [anhydrous ethanol R](#) and dilute to 100.0 mL with the same solvent.

Test solution Dissolve 0.200 g of the substance to be examined in 25.0 mL of the internal standard solution. Add 25 mL of [water R](#) and shake with 4 quantities, each of 25 mL, of [pentane R](#), adding [sodium chloride R](#), if necessary, to facilitate the separation of the layers. Combine the upper layers, wash with 2 quantities, each of 30 mL, of [water R](#), dry over [anhydrous sodium sulfate R](#) and filter.

Reference solution (a) Dissolve 0.100 g of [cetyl alcohol CRS](#) in 25.0 mL of the internal standard solution. Add 25 mL of [water R](#) and shake with 4 quantities, each of 25 mL, of [pentane R](#), adding [sodium chloride R](#), if necessary, to facilitate the separation of the layers. Combine the upper layers, wash with 2 quantities, each of 30 mL, of [water R](#), dry over [anhydrous sodium sulfate R](#) and filter.

Reference solution (b) Dissolve 0.100 g of [stearyl alcohol CRS](#) in 25.0 mL of the internal standard solution. Add 25 mL of [water R](#) and shake with 4 quantities, each of 25 mL, of [pentane R](#), adding [sodium chloride R](#), if necessary, to facilitate the separation of the layers. Combine the upper layers, wash with 2 quantities, each of 30 mL, of [water R](#), dry over [anhydrous sodium sulfate R](#) and filter.

Column:

- *material:* fused silica;
- *size:* $l = 25\text{ m}$, $\varnothing = 0.25\text{ mm}$;
- *stationary phase:* [methylpolysiloxane R](#) (film thickness 0.25 μm).

Carrier gas [helium for chromatography R](#).

Flow rate 1 mL/min.

Split ratio 1:100.

Temperature:

	Time (min)	Temperature (°C)
Column	0 - 20	150 → 250
Injection port		250
Detector		250

Detection Flame ionisation.

Injection 1 μL .

Elution order Cetyl alcohol, stearyl alcohol, 1-nonadecanol.

Calculate the percentage content of cetyl alcohol in the substance to be examined using the following expression and taking into account the assigned content of [cetyl alcohol CRS](#):

— — —

- A_x = area of the peak due to cetyl alcohol in the chromatogram obtained with the test solution;
- $A_{x,y}$ = area of the peak due to [cetyl alcohol CRS](#) in the chromatogram obtained with reference solution (a);
- A_1 = area of the peak due to the internal standard in the chromatogram obtained with the test solution;
- A_2 = area of the peak due to the internal standard in the chromatogram obtained with reference solution (a);
- m = mass of the substance to be examined in the test solution, in milligrams;

$m_{x,y}$ = mass of [cetyl alcohol CRS](#) in reference solution (a), in milligrams.

Calculate the percentage content of stearyl alcohol in the substance to be examined using the following expression and taking into account the assigned content of [stearyl alcohol CRS](#):

— — —

A_z = area of the peak due to stearyl alcohol in the chromatogram obtained with the test solution;

$A_{z,y}$ = area of the peak due to [stearyl alcohol CRS](#) in the chromatogram obtained with reference solution (b);

A_1 = area of the peak due to the internal standard in the chromatogram obtained with the test solution;

A_3 = area of the peak due to the internal standard in the chromatogram obtained with reference solution (b);

m = mass of the substance to be examined in the test solution, in milligrams;

$m_{z,y}$ = mass of [stearyl alcohol CRS](#) in reference solution (b), in milligrams.

The percentage content of cetostearyl alcohol corresponds to the sum of the percentage contents of cetyl alcohol and stearyl alcohol.

[Sodium laurilsulfate](#)

Disperse 0.300 g in 25 mL of [methylene chloride R](#). Add 50 mL of [water R](#) and 10 mL of [dimidium bromide-sulfan blue mixed solution R](#). Titrate with [0.004 M benzethonium chloride](#), using sonication, heating, and allowing the layers to separate before each addition, until the colour of the lower layer changes from pink to grey.

1 mL of [0.004 M benzethonium chloride](#) is equivalent to 1.154 mg of sodium laurilsulfate.

LABELLING

The label states, where applicable, the name and concentration of any added buffer.

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