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Macrogols

[General Notices](#)

(Ph. Eur. monograph 1444)

Action and use

Non-ionic surfactant.

Ph Eur

DEFINITION

Mixtures of polymers with the general formula $H-[OCH_2-CH_2]_n-OH$ where n represents the average number of oxyethylene groups. The type of macrogol is defined by a number that indicates the average relative molecular mass. A suitable stabiliser may be added.

CHARACTERS

Type of macrogol	Appearance	Solubility
300	clear, viscous, colourless or almost	miscible with water, very soluble in acetone, in ethanol (96 per cent) and in methylene chloride, practically insoluble in fatty oils and in mineral oils
400	colourless, hygroscopic liquid	
600		
1000	white or almost white, hygroscopic solid with a waxy or paraffin-like appearance	very soluble in water, freely soluble in ethanol (96 per cent) and in methylene chloride, practically insoluble in fatty oils and in mineral oils
1500	white or almost white solid with a waxy or paraffin-like appearance	very soluble in water and in methylene chloride, freely soluble in ethanol (96 per cent), practically insoluble in fatty oils and in mineral oils
3000	white or almost white solid with a waxy or paraffin-like appearance	very soluble in water and in methylene chloride, very slightly soluble in ethanol (96 per cent), practically insoluble in fatty oils and in mineral oils
3350		
4000	white or almost white solid with a waxy or paraffin-like appearance	very soluble in water and in methylene chloride, practically insoluble in ethanol (96 per cent), in fatty oils and in mineral oils
6000		
8000		

Type of macrogol	Appearance	Solubility
20 000	white or almost white solid with a waxy or	very soluble in water, soluble in methylene chloride, practically
35 000	paraffin-like appearance	insoluble in ethanol (96 per cent), in fatty oils and in mineral oils

IDENTIFICATION

A. Viscosity (see Tests).

B. To 1 g in a test-tube add 0.5 mL of [sulfuric acid R](#), close the test-tube with a stopper fitted with a bent delivery tube and heat until white fumes are evolved. Collect the fumes via the delivery tube into 1 mL of [mercuric chloride solution R](#). An abundant, white, crystalline precipitate is formed.

C. To 0.1 g add 0.1 g of [potassium thiocyanate R](#) and 0.1 g of [cobalt nitrate R](#) and mix thoroughly with a glass rod. Add 5 mL of [methylene chloride R](#) and shake. The liquid phase becomes blue.

TESTS

Appearance of solution

The solution is clear ([2.2.1](#)) and not more intensely coloured than reference solution BY₆ ([2.2.2, Method II](#)).

Dissolve 12.5 g in [water R](#) and dilute to 50 mL with the same solvent.

Acidity or alkalinity

Dissolve 5.0 g in 50 mL of [carbon dioxide-free water R](#) and add 0.15 mL of [bromothymol blue solution R1](#). The solution is yellow or green. Not more than 0.1 mL of [0.1 M sodium hydroxide](#) is required to change the colour of the indicator to blue.

Viscosity ([2.2.9](#))

The viscosity is calculated using a density given in Table 1444.-1.

For macrogols with a relative molecular mass greater than 400, determine the viscosity on a 50 per cent *m/m* solution of the substance to be examined.

Freezing point ([2.2.18](#))

See Table 1444.-2.

Hydroxyl value

Introduce *m* g (see Table 1444.-3) into a dry conical flask fitted with a reflux condenser. Add 25.0 mL of [phthalic anhydride solution R](#), swirl to dissolve and boil under a reflux condenser on a hot plate for 60 min. Allow to cool. Rinse the condenser first with 25 mL of [pyridine R](#) and then with 25 mL of [water R](#), add 1.5 mL of [phenolphthalein solution R](#) and titrate with [1 M sodium hydroxide](#) until a faint pink colour is obtained (*n*₁ mL). Carry out a blank test (*n*₂ mL). Calculate the hydroxyl value using the following expression:

$$\frac{56.1 \times (n_2 - n_1)}{m}$$

For macrogols with a relative molecular mass greater than 1000, if the water content is more than 0.5 per cent, dry a sample of suitable mass at 100-105 °C for 2 h and carry out the determination of the hydroxyl value on the dried sample.

Table 1444.-1

Type of macrogol	Kinematic viscosity (mm ² ·s ⁻¹)	Dynamic viscosity (mPa·s)	Density* (g/mL)
300	71 - 94	80 - 105	1.120
400	94 - 116	105 - 130	1.120
600	13.9 - 18.5	15 - 20	1.080
1000	20.4 - 27.7	22 - 30	1.080
1500	31 - 46	34 - 50	1.080
3000	69 - 93	75 - 100	1.080
3350	76 - 110	83 - 120	1.080
4000	102 - 158	110 - 170	1.080
6000	185 - 250	200 - 270	1.080
8000	240 - 472	260 - 510	1.080
20 000	2500 - 3200	2700 - 3500	1.080
35 000	10 000 - 13 000	11 000 - 14 000	1.080

*Density of the substance for macrogols 300 and 400. Density of the 50 per cent *m/m* solution for the other macrogols.

Table 1444.-2

Type of macrogol	Freezing point (°C)
600	15 - 25
1000	35 - 40
1500	42 - 48
3000	50 - 56
3350	53 - 57

Type of macrogol**Freezing point****(°C)**

4000	53 - 59
6000	55 - 61
8000	55 - 62
20 000	minimum 57
35 000	minimum 57

Table 1444.-3

Type of macrogol	Hydroxyl value	m (g)
300	340 - 394	1.5
400	264 - 300	1.9
600	178 - 197	3.5
1000	107 - 118	5.0
1500	70 - 80	7.0
3000	34 - 42	12.0
3350	30 - 38	12.0
4000	25 - 32	14.0
6000	16 - 22	18.0
8000	12 - 16	24.0
20 000	-	-
35 000	-	-

Reducing substances

Dissolve 1 g in 1 mL of a 10 g/L solution of [resorcinol R](#) and warm gently if necessary. Add 2 mL of [hydrochloric acid R](#). After 5 min the solution is not more intensely coloured than reference solution R₃ ([2.2.2, Method I](#)).

[Formaldehyde](#)

Maximum 30 ppm.

Test solution To 1.00 g add 0.25 mL of [chromotropic acid, sodium salt solution R](#), cool in iced water and add 5.0 mL of [sulfuric acid R](#). Allow to stand for 15 min and dilute slowly to 10 mL with [water R](#).

Reference solution Dilute 0.860 g of [formaldehyde solution R](#) to 100 mL with [water R](#). Dilute 1.0 mL of this solution to 100 mL with [water R](#). In a 10 mL flask, mix 1.00 mL of this solution and 0.25 mL of [chromotropic acid, sodium salt solution R](#), cool in iced water and add 5.0 mL of [sulfuric acid R](#). Allow to stand for 15 min and dilute slowly to 10 mL with [water R](#).

Blank solution In a 10 mL flask mix 1.00 mL of [water R](#) and 0.25 mL of [chromotropic acid, sodium salt solution R](#), cool in iced water and add 5.0 mL of [sulfuric acid R](#). Dilute slowly to 10 mL with [water R](#).

Determine the absorbance ([2.2.25](#)) of the test solution at 567 nm, against the blank solution. It is not higher than that of the reference solution.

If the use of macrogols with a higher content of formaldehyde may have adverse effects, the competent authority may impose a limit of not more than 15 ppm.

Ethylene glycol and diethylene glycol

carry out this test only if the macrogol has a relative molecular mass below 1000.

Gas chromatography ([2.2.28](#)).

Test solution Dissolve 5.00 g of the substance to be examined in [acetone R](#) and dilute to 100.0 mL with the same solvent.

Reference solution Dissolve 0.10 g of [ethylene glycol R](#) and 0.50 g of [diethylene glycol R](#) in [acetone R](#) and dilute to 100.0 mL with the same solvent. Dilute 1.0 mL of the solution to 10.0 mL with [acetone R](#).

Column:

- *material:* glass;
- *size:* $l = 1.8 \text{ m}$, $\varnothing = 2 \text{ mm}$;
- *stationary phase:* [silanised diatomaceous earth for gas chromatography R](#), impregnated with 5 per cent *m/m* of [macrogol 20 000 R](#).

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

Flow rate 30 mL/min.

Temperature:

- *column:* if necessary, precondition the column by heating at 200 °C for about 15 h; adjust the initial temperature of the column to obtain a retention time of 14-16 min for diethylene glycol; raise the temperature of the column by about 30 °C at a rate of 2 °C/min but without exceeding 170 °C;
- *injection port and detector:* 250 °C.

Detection Flame ionisation.

Injection 2 µL.

Carry out 5 replicate injections to check the repeatability of the response.

Limit Maximum 0.4 per cent, calculated as the sum of the contents of ethylene glycol and diethylene glycol.

[Ethylene oxide and dioxan \(2.4.25\)](#)

Maximum 1 ppm of ethylene oxide and 10 ppm of dioxan.

[Water \(2.5.12\)](#)

Maximum 2.0 per cent for macrogols with a relative molecular mass not greater than 1000 and maximum 1.0 per cent for macrogols with a relative molecular mass greater than 1000, determined on 2.00 g.

[Sulfated ash \(2.4.14\)](#)

Maximum 0.2 per cent, determined on 1.0 g.

STORAGE

In an airtight container.

LABELLING

The label states:

- the type of macrogol;
- the content of formaldehyde.

FUNCTIONALITY-RELATED CHARACTERISTICS

This section provides information on characteristics that are recognised as being relevant control parameters for one or more functions of the substance when used as an excipient (see chapter 5.15). Some of the characteristics described in the Functionality-related characteristics section may also be present in the mandatory part of the monograph since they also represent mandatory quality criteria. In such cases, a cross-reference to the tests described in the mandatory part is included in the Functionality-related characteristics section. Control of the characteristics can contribute to the quality of a medicinal product by improving the consistency of the manufacturing process and the performance of the medicinal product during use. Where control methods are cited, they are recognised as being suitable for the purpose, but other methods can also be used. Wherever results for a particular characteristic are reported, the control method must be indicated.

The following characteristic may be relevant for macrogols used as solvent.

[Viscosity](#)

(see Tests).

The following characteristics may be relevant for macrogols used as suspension stabiliser and thickener.

[Viscosity](#)

(see Tests).

The following characteristic may be relevant for macrogols used as lubricant in tablets.

Particle-size distribution ([2.9.31](#))

The following characteristics may be relevant for macrogols used as suppository base and for macrogols used in hydrophilic ointments.

[Viscosity](#)

(see Tests).

Melting point ([2.2.15](#))

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