Containers and storage Containers—Tight containers.
Storage—Light-resistant.

Betamethasone Dipropionate
ジプロピオン酸ベタメタゾン

\[
\text{C}_{28}\text{H}_{32}\text{FO}_{7}: \text{504.59}
\]

9-Fluoro-11\beta,17,21-trihydroxy-16\beta-methylpregna-1,4-
diene-3,20-dione 17,21-dipropionate [5593-20-4]

Betamethasone Dipropionate, when dried, contains not less than 97.0% and not more than 103.0% of \( \text{C}_{28}\text{H}_{32}\text{FO}_{7} \), and not less than 3.4% and not more than 4.1% of fluorine (F:19.00).

Description Betamethasone Dipropionate occurs as a white to pale yellowish white, crystalline powder. It is odorless.

It is freely soluble in acetone, in 1,4-dioxane and in chloroform, soluble in methanol, sparingly soluble in ethanol (95), slightly soluble in diethyl ether, and practically insoluble in water and in hexane.

It is affected gradually by light.

Identification (1) To 1 mL of a solution of Betamethasone Dipropionate in methanol (1 in 10,000) add 4 mL of isonicotinic acid TS, and heat on a water bath for 2 minutes: a yellow color develops.

(2) Proceed with 0.01 g of Betamethasone Dipropionate as directed under the Oxygen Flask Combustion Method, using a mixture of 0.5 mL of 0.01 mol/L sodium hydroxide TS and 20 mL of water as the absorbing liquid: the test solution so obtained responds to the Qualitative Tests for fluoride.

(3) Determine the absorption spectrum of a solution of Betamethasone Dipropionate in methanol (3 in 200,000) as directed under the Ultraviolet-visible Spectrophotometry, and compare the spectrum with the Reference Spectrum: both spectra exhibit similar intensities of absorption at the same wavelengths.

(4) Determine the infrared absorption spectrum of Betamethasone Dipropionate, previously dried, as directed in the potassium bromide disk method under the Infrared Spectrophotometry, and compare the spectrum with the Reference Spectrum: both spectra exhibit similar intensities of absorption at the same wave numbers.

Melting point 176 - 180°C

Optical rotation \([\alpha]_{D}^{20}: +63 - +70^{\circ}\) (after drying, 0.05 g, 1,4-dioxane, 10 mL, 100 mm).

Purity (1) Fluoride—To 0.10 g of Betamethasone Dipropionate add 10.0 mL of diluted 0.01 mol/L sodium hydroxide TS (1 in 20), shake for 10 minutes, and filter through a membrane filter (0.4-μm pore size). Place 5.0 mL of the filtrate in a 20-mL volumetric flask, and add 10 mL of a mixture of alizarin complexone TS, acetic acid-potassium acetate buffer solution, pH 4.3, and cerium (III) nitrate TS (1:1:1), add water to make 20 mL, allow to stand for 1 hour, and use this solution as the sample solution. Separately, place 1.0 mL of Standard Fluorine Solution in a 20-mL volumetric flask, add 5.0 mL of diluted 0.01 mol/L sodium hydroxide TS (1 in 20), then 10 mL of a mixture of alizarin complexone TS, acetic acid-potassium acetate buffer solution, pH 4.3, and cerium (III) nitrate TS (1:1:1), proceed in the same manner as the preparation of the sample solution, and use this solution as the standard solution. Place 5.0 mL of diluted 0.01 mol/L sodium hydroxide TS (1 in 20) in a 20-mL volumetric flask, and proceed in the same manner as the preparation of the sample solution. Using this solution as the blank, determine the absorbances of the sample solution and the standard solution at 600 nm as directed under the Ultraviolet-visible Spectrophotometry: the absorbance of the sample solution is not greater than that of the standard solution (not more than 0.012%).

(2) Heavy metals—Proceed with 1.0 g of Betamethasone Dipropionate according to Method 2, and perform the test. Prepare the control solution with 2.0 mL of Standard Lead Solution (not more than 20 ppm).

(3) Other steroids—Conduct this procedure without exposure to daylight, using light-resistant vessels. Dissolve 0.010 g of Betamethasone Dipropionate in 10 mL of chloroform, and use this solution as the sample solution. Pipet 3 mL of the sample solution, add chloroform to make exactly 100 mL, and use this solution as the standard solution. Perform the test as directed under the Thin-layer Chromatography with these solutions. Spot 20 \( \mu \)L each of the sample solution and the standard solution on a plate of silica gel with fluorescent indicator for thin-layer chromatography. Develop the plate with a mixture of chloroform and acetone (7:1) to a distance of about 10 cm, and air-dry the plate. Examine under ultraviolet light (main wavelength: 254 nm): the spots other than the principal spot from the sample solution are not more intense than the spot from the standard solution.

Loss on drying Not more than 1.0% (0.5 g, 105°C, 3 hours).

Residue on ignition Not more than 0.2% (0.5 g, platinum crucible).

Assay (1) Betamethasone dipropionate—Weigh accurately about 0.015 g of Betamethasone Dipropionate, previously dried, and dissolve in methanol to make exactly 100 mL. Pipet 5 mL of this solution, and dilute with methanol to exactly 50 mL. Determine the absorbance \( A \) of this solution at the wavelength of maximum absorption at about 239 nm as directed under the Ultraviolet-visible Spectrophotometry.

\[
\text{Amount (mg)} \ of \ \text{C}_{28}\text{H}_{32}\text{FO}_{7} = \frac{A}{312} \times 10,000
\]

(2) Fluorine—Weigh accurately about 0.01 g of Betamethasone Dipropionate, previously dried, and proceed as directed in the procedure of determination for fluorine under the Oxygen Flask Combustion Method, using a mixture
of 0.5 mL of 0.01 mol/L sodium hydroxide TS and 20 mL of water as the absorbing liquid.

Containers and storage Containers—Tight containers.
Storage—Light-resistant.

Betamethasone Sodium Phosphate

リン酸ベタメタゾントリウム

\[ \text{C}_{29}\text{H}_{38}\text{FNa}_{2}\text{O}_{3}\text{P} : 516.40} \]

Disodium 9-fluoro-11β,17,21-trihydroxy-16β-
methylpregna-1,4-diene-3,20-dione 21-phosphate

[151-73-5]

Betamethasone Sodium Phosphate contains not less than
97.0% and not more than 103.0% of
\[ \text{C}_{29}\text{H}_{38}\text{FNa}_{2}\text{O}_{3}\text{P} \], calculated on the anhydrous basis.

Description Betamethasone Sodium Phosphate occurs as
white to pale yellowish white, crystalline powder or masses.
It is odorless.
It is freely soluble in water, sparingly soluble in methanol,
slightly soluble in ethanol (95), and practically insoluble in
diethyl ether.
It is hygroscopic.
Meltng point: about 213°C (with decomposition).

Identification (1) Dissolve 2 mg of Betamethasone Sodium
Phosphate in 2 mL of sulfuric acid: a brown color de-
velops, and gradually changes to blackish brown.
(2) Prepare the test solution with 0.01 g of Betamethasone
Sodium Phosphate as directed under the Oxygen Flask
Combustion Method, using a mixture of 0.5 mL of 0.01
mol/L sodium hydroxide TS and 20 mL of water as an ab-
sorbing liquid: the test solution responds to the Qualitative
Tests (2) for fluoride.
(3) Take 0.04 g of Betamethasone Sodium Phosphate in
a platinum crucible, and carbonize by heating. After cool-
ing, add 5 drops of nitric acid, and incinerate by heating.
To the residue add 10 mL of diluted nitric acid (1 in 50),
and boil for several minutes. After cooling, neutralize the
solution with ammonia TS, filter if necessary, and use this solu-
tion as the sample solution. The sample solution responds to the Qualitative Tests for sodium salt and for phosphate.
(4) Determine the infrared absorption spectrum of Be-
tamethasone Sodium Phosphate, previously dried, as direc-
ted in the potassium bromide disk method under the Infrared
Spectrophotometry, and compare the spectrum with the
Reference Spectrum or the spectrum of previously dried Be-
tamethasone Sodium Phosphate Reference Standard: both spectra exhibit similar intensities of absorption at the same
wave numbers.

Optical rotation \[ [\alpha]_D^{20} +99 +105^\circ \] (0.1 g, calculated on the anhydrous basis, water, 10 mL, 100 mm).

pH Dissolve 0.10 g of Betamethasone Sodium Phosphate
in 20 mL of water: the pH of this solution is between 7.5 and
9.0.

Purity (1) Clarity and color of solution—Dissolve 0.25 g of
Betamethasone Sodium Phosphate in 10 mL of water: the
solution is clear and colorless.
(2) Free phosphoric acid—Weigh accurately about 0.02 g
of Betamethasone Sodium Phosphate, dissolve in 20 mL of
water, and use this solution as the sample solution. Sepa-
rately, pipet 4 mL of Standard Phosphoric Acid Solu-
tion, add 20 mL of water, and use this solution as the stan-
dard solution. To each of the sample solution and the stan-
dard solution add exactly 7 mL of dilute sulfuric acid, exact-
ly 2 mL of hexaammonium heptamolybdate-sulfuric acid
TS and exactly 2 mL of \( p \)-methylaminophenol sulfonate TS,
shake well, and allow to stand at 20 ± 1°C for 15 minutes.
To each add water to make exactly 50 mL, and allow to
stand at 20 ± 1°C for 15 minutes. Perform the test with
these solutions as directed under the Ultraviolet-visible Spec-
trophotometry, using a solution prepared with 20 mL of
water in the same manner as the blank. Determine the absorb-
cances, \( \lambda \) and \( A_s \), of each solution from the sample
solution and the standard solution at 730 nm: the amount of
free phosphoric acid is not more than 0.5%.

\[ \text{Amount (\%)} \times \frac{\lambda}{A_s} \times 10.32 \]

\( W \): Amount (mg) of Betamethasone Sodium Phosphate,
calculated on the anhydrous basis.

(3) Betamethasone—Dissolve 0.020 g of Betamethasone
Sodium Phosphate in exactly 2 mL of methanol, and use
this solution as the sample solution. Separately, dissolve
0.020 g of Betamethasone Reference Standard in exactly 10
mL of methanol. Pipet 1 mL of this solution, add methanol
to make exactly 20 mL, and use this solution as the standard
solution. Perform the test with these solutions as directed
under the Thin-layer Chromatography. Spot 5 \( \mu \)L each of
the sample solution and the standard solution on a plate of
silica gel with fluorescent indicator for thin-layer chroma-
tography. Develop the plate with a freshly prepared mix-
ture of 1-butanol, water and acetic anhydride (3:1:1) to a dis-
tance of about 10 cm, and air-dry the plate. Examine under
ultraviolet light (main wavelength: 254 nm): the spot from
the sample solution corresponding to the spot from the stan-
dard solution is not more intense than the spot from the stan-
dard solution.

Water Not more than 10.0% (0.2 g, back titration).

Assay Weigh accurately about 0.02 g each of Betama-
thesone Sodium Phosphate and Betamethasone Sodium Phos-
phate Reference Standard (determine its water content be-
fore using in the same manner as Betamethasone Sodium
Phosphate), and dissolve each in methanol to make exactly
20 mL. Pipet 5 mL each of these solutions, add exactly 5
mL of the internal standard solution, then add methanol
to make 50 mL, and use these solutions as the sample solution
and the standard solution, respectively. Perform the test
with 10 \( \mu \)L each of the sample solution and the standard so-
lution as directed under the Liquid Chromatography accord-
ing to the following conditions, and calculate the ratios, \( Q_T \)
and \( Q_s \), of the peak area of betamethasone phosphate to
that of the internal standard, respectively.