

Status: Currently Official on 14-Feb-2025  
 Official Date: Official as of 01-May-2016  
 Document Type: USP Monographs  
 DocId: GUID-1E695F8E-1201-46A4-9B37-ADC02D7A1364\_2\_en-US  
 DOI: [https://doi.org/10.31003/USPNF\\_M23510\\_02\\_01](https://doi.org/10.31003/USPNF_M23510_02_01)  
 DOI Ref: 56g0y

© 2025 USPC  
 Do not distribute

## Dexamethasone Sodium Phosphate Ophthalmic Solution

### DEFINITION

Dexamethasone Sodium Phosphate Ophthalmic Solution is a sterile, aqueous solution of Dexamethasone Sodium Phosphate. It contains an amount of dexamethasone sodium phosphate ( $C_{22}H_{28}FNa_2O_8P$ ) equivalent to NLT 90.0% and NMT 115.0% of the labeled amount of dexamethasone phosphate ( $C_{22}H_{30}FO_8P$ ).

### IDENTIFICATION

#### • A. THIN-LAYER CHROMATOGRAPHY

**Solution A:** Dissolve 3.1 g of boric acid, 203 mg of magnesium chloride, and 860 mg of sodium hydroxide in enough water to make 1000 mL.

**Solution B:** 1 mg/mL of alkaline phosphatase enzyme in *Solution A*

**Standard solution:** 300 µg/mL of [USP Dexamethasone RS](#) in methylene chloride

**Sample solution:** Transfer 5 mL of *Solution B* to a glass-stoppered, 50-mL tube containing 5 mL of the *Sample solution* from the Assay.

Incubate at 37° for 45 min, then add 25 mL of methylene chloride and shake for 2 min. Evaporate 15 mL of the methylene chloride extract on a steam bath to dryness, and dissolve the residue in 1 mL of methylene chloride.

#### Chromatographic system

(See [Chromatography \(621\)](#), [Thin-Layer Chromatography](#).)

**Adsorbent:** 0.25-mm layer of chromatographic silica gel mixture (20- × 20-cm plate)

**Application volume:** 5 µL

**Developing solvent system:** Chloroform, acetone, and water (50:50:1)

**Spray reagent:** Dilute sulfuric acid (1 in 2)

#### Analysis

**Samples:** *Standard solution* and *Sample solution*

Allow the spots to dry, and develop the chromatogram using the *Developing solvent system* in a tank completely lined with filter paper until the solvent front has moved about three-fourths of the length of the plate. Remove the plate from the developing tank, mark the solvent front, and allow the spots to dry. Spray the plate with *Spray reagent*, and heat at 105° until brown or black spots appear.

**Acceptance criteria:** The  $R_f$  value of the principal spot of the *Sample solution* corresponds to that of the *Standard solution*.

• **B.** The retention time of the major peak of the *Sample solution* corresponds to that of the *Standard solution*, as obtained in the Assay.

### ASSAY

#### • PROCEDURE

**Mobile phase:** 0.01 M monobasic potassium phosphate in a mixture of methanol and water (1:1)

**Standard solution:** 0.09 mg/mL of freshly prepared [USP Dexamethasone Sodium Phosphate RS](#) in *Mobile phase*

**Sample solution:** Nominally 0.08 mg/mL of dexamethasone phosphate from Ophthalmic Solution in *Mobile phase*

#### Chromatographic system

(See [Chromatography \(621\)](#), [System Suitability](#).)

**Mode:** LC

**Detector:** UV 254 nm

**Column:** 4-mm × 30-cm; packing L1

**Flow rate:** 1.6 mL/min

**Injection volume:** 20 µL

#### System suitability

**Sample:** *Standard solution*

[NOTE—The retention time for dexamethasone phosphate is about 5 min.]

#### Suitability requirements

**Relative standard deviation:** NMT 1.5%

#### Analysis

**Samples:** *Standard solution* and *Sample solution*

Calculate the percentage of the labeled amount of dexamethasone phosphate ( $C_{22}H_{30}FO_8P$ ) in the portion of Ophthalmic Solution taken:

$$\text{Result} = (r_U/r_S) \times (C_S/C_U) \times (M_{r1}/M_{r2}) \times 100$$

$r_U$  = peak response from the *Sample solution*

$r_S$  = peak response from the *Standard solution*

$C_S$  = concentration of [USP Dexamethasone Sodium Phosphate RS](#) in the *Standard solution* (mg/mL)

$C_U$  = nominal concentration of dexamethasone phosphate in the *Sample solution* (mg/mL)

$M_{r1}$  = molecular weight of dexamethasone phosphate, 472.44

$M_{r2}$  = molecular weight of dexamethasone sodium phosphate, 516.40

**Acceptance criteria:** 90.0%–115.0%

#### SPECIFIC TESTS

- **pH** (791): 6.6–7.8
- **STERILITY TESTS** (71): Meets the requirements

#### ADDITIONAL REQUIREMENTS

- **PACKAGING AND STORAGE:** Preserve in tight, light-resistant containers. Store between 15° and 25°.
- **USP REFERENCE STANDARDS** (11).  
[USP Dexamethasone RS](#)  
[USP Dexamethasone Sodium Phosphate RS](#)

**Auxiliary Information** - Please [check for your question in the FAQs](#) before contacting USP.

Topic/Question	Contact	Expert Committee
DEXAMETHASONE SODIUM PHOSPHATE OPHTHALMIC SOLUTION	<a href="#">Documentary Standards Support</a>	SM52020 Small Molecules 5

**Chromatographic Database Information:** [Chromatographic Database](#)

#### Most Recently Appeared In:

Pharmacopeial Forum: Volume No. PF 43(6)

**Current DocID:** GUID-1E695F8E-1201-46A4-9B37-ADC02D7A1364\_2\_en-US

**Previous DocID:** GUID-1E695F8E-1201-46A4-9B37-ADC02D7A1364\_1\_en-US

**DOI:** [https://doi.org/10.31003/USPNF\\_M23510\\_02\\_01](https://doi.org/10.31003/USPNF_M23510_02_01)

**DOI ref:** [56g0y](#)