Status: Currently Official on 14-Feb-2025
Official Date: Official as of 01-May-2020
Document Type: USP Monographs
DocId: GUID-BAA1004B-2261-4FEA-9E4B-058A8F56DFC6_5_en-US
DOI: https://doi.org/10.31003/USPNF_M29350_05_01
DOI Ref: x6eql

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Ephedrine Hydrochloride

C₁₀H₁₅NO · HCI

201.69

Benzenemethanol, α -[1-(methylamino)ethyl]-, hydrochloride, [R-(R*,S*)]-;

(-)-Ephedrine hydrochloride;

(1R,2S)-2-(Methylamino)-1-phenylpropan-1-ol hydrochloride; CAS RN®: 50-98-6; UNII: NLJ6390P1Z.

DEFINITION

Ephedrine Hydrochloride contains NLT 98.0% and NMT 102.0% of ephedrine hydrochloride (C₁₀H₁₅NO·HCl), calculated on the dried basis.

IDENTIFICATION

Change to read:

• A. <u>Spectroscopic Identification Tests (197), Infrared Spectroscopy</u>: 197K_{▲ (CN 1-May-2020)}

• B. IDENTIFICATION TESTS—GENERAL (191), Chemical Identification Tests, Chloride: Meets the requirements

• C. The retention time of the ephedrine peak of the Sample solution corresponds to that of the Standard solution, as obtained in the Assay.

ASSAY

• PROCEDURE

Buffer: 11.6 g/L of ammonium acetate. Adjust with glacial acetic acid to a pH of 4.0.

Mobile phase: Methanol and Buffer (6:94) **Diluent:** Methanol and water (6:94)

System suitability solution: 0.1 mg/mL each of USP Ephedrine Hydrochloride RS and USP Pseudoephedrine Hydrochloride RS in Diluent

Standard solution: 0.2 mg/mL of <u>USP Ephedrine Hydrochloride RS</u> in *Diluent*

Sample solution: 0.2 mg/mL of Ephedrine Hydrochloride in *Diluent*

Chromatographic system

(See Chromatography (621), System Suitability.)

Mode: LC

Detector: UV 257 nm

Column: 4.6-mm × 15-cm; 3-µm packing L11

Flow rate: 1 mL/min Injection volume: 20 μL

Run time: NLT 2 times the retention time of ephedrine

System suitability

Samples: System suitability solution and Standard solution

[Note—The relative retention times for ephedrine and pseudoephedrine are 1.0 and 1.1, respectively.]

Suitability requirements

Resolution: NLT 2.0 between ephedrine and pseudoephedrine, System suitability solution

Tailing factor: NMT 2.0 for ephedrine, Standard solution

Relative standard deviation: NMT 0.73% for ephedrine, Standard solution

Analysis

Samples: Standard solution and Sample solution

Calculate the percentage of ephedrine hydrochloride ($C_{10}H_{15}NO \cdot HCI$) in the portion of Ephedrine Hydrochloride taken:

Result =
$$(r_{II}/r_{S}) \times (C_{S}/C_{II}) \times 100$$

r,, = peak response of ephedrine from the Sample solution

 $r_{\rm s}$ = peak response of ephedrine from the Standard solution

C_s = concentration of <u>USP Ephedrine Hydrochloride RS</u> in the *Standard solution* (mg/mL)

C₁₁ = concentration of Ephedrine Hydrochloride in the Sample solution (mg/mL)

Acceptance criteria: 98.0%-102.0% on the dried basis

IMPURITIES

• Residue on Ignition (281): NMT 0.1%

• CHLORIDE AND SULFATE (221), Sulfate

Sample solution: 1.25 mg/mL of Ephedrine Hydrochloride in water

Analysis: Add 1 mL of 3 N hydrochloric acid and 1 mL of barium chloride TS to 40 mL of the Sample solution.

Acceptance criteria: No turbidity develops within 10 min.

• ORGANIC IMPURITIES

Buffer, Mobile phase, and Chromatographic system: Proceed as directed in the Assay, except for the Run time.

Run time: NLT 2.5 times the retention time of ephedrine

System suitability solution: 0.1 mg/mL each of USP Ephedrine Hydrochloride RS and USP Pseudoephedrine Hydrochloride RS in Mobile

phase

Sensitivity solution: $3.8 \mu g/mL$ of <u>USP Ephedrine Hydrochloride RS</u> in *Mobile phase* Standard solution: $30 \mu g/mL$ of <u>USP Ephedrine Hydrochloride RS</u> in *Mobile phase*

Sample solution: 7.5 mg/mL of Ephedrine Hydrochloride in Mobile phase

System suitability

Samples: System suitability solution, Sensitivity solution, and Standard solution

[Note—See <u>Table 1</u> for the relative retention times.]

Suitability requirements

Resolution: NLT 2.0 between ephedrine and pseudoephedrine, System suitability solution

Relative standard deviation: NMT 5.0%, Standard solution Signal-to-noise ratio: NLT 10, Sensitivity solution

Analysis

Samples: Standard solution and Sample solution

Calculate the percentage of α -acetylbenzyl alcohol or any unspecified impurity in the portion of Ephedrine Hydrochloride taken:

Result =
$$(r_{ij}/r_{s}) \times (C_{s}/C_{ij}) \times (1/F) \times 100$$

 r_{ij} = peak response of α -acetylbenzyl alcohol or any unspecified impurity from the Sample solution

 $r_{\rm s}$ = peak response of ephedrine from the Standard solution

C_s = concentration of <u>USP Ephedrine Hydrochloride RS</u> in the *Standard solution* (mg/mL)

C₁₁ = concentration of Ephedrine Hydrochloride in the Sample solution (mg/mL)

F = relative response factor (see <u>Table 1</u>)

Acceptance criteria: See <u>Table 1</u>. The reporting threshold is 0.05%.

Table 1

Name	Relative Retention Time	Relative Response Factor	Acceptance Criteria, NMT (%)
Ephedrine	1.0	_	-
Pseudoephedrine ^a	1.1	_	-
α-Acetylbenzyl alcohol ^{<u>b</u>}	1.4	2.5	0.2
Any unspecified impurity	-	1.0	0.1
Total impurities [©]	-	_	0.5

^a Included for identification only. It is not to be reported and not to be included in the total impurities.

b (-)-(1*R*)-1-Hydroxy-1-phenylpropan-2-one.

https://trumgtamthuoc.com/

 $^{\text{c}}$ Excludes α -acetylbenzyl alcohol.

SPECIFIC TESTS

• OPTICAL ROTATION (781S), Procedures, Specific Rotation

Sample solution: 50 mg/mL of Ephedrine Hydrochloride in water

Acceptance criteria: -33.0° to -35.5°

• Loss on Drying (731)

Analysis: Dry at 105° for 3 h. **Acceptance criteria:** NMT 0.5%

• ACIDITY OR ALKALINITY

Sample solution: 50 mg/mL of Ephedrine Hydrochloride in water **Analysis:** To 20 mL of *Sample solution* add 1 drop of methyl red TS.

Acceptance criteria: If the solution is yellow, it is changed to red by NMT 0.10 mL of 0.020 N sulfuric acid. If the solution is pink, it is changed

to yellow by NMT 0.20 mL of 0.020 N sodium hydroxide.

ADDITIONAL REQUIREMENTS

• Packaging and Storage: Preserve in well-closed, light-resistant containers.

USP REFERENCE STANDARDS (11)
 USP Ephedrine Hydrochloride RS
 USP Pseudoephedrine Hydrochloride RS

Auxiliary Information - Please check for your question in the FAQs before contacting USP.

Topic/Question	Contact	Expert Committee
EPHEDRINE HYDROCHLORIDE	Documentary Standards Support	SM52020 Small Molecules 5

Chromatographic Database Information: Chromatographic Database

Most Recently Appeared In:

Pharmacopeial Forum: Volume No. PF 44(2)

Current DocID: GUID-BAA1004B-2261-4FEA-9E4B-058A8F56DFC6_5_en-US

DOI: https://doi.org/10.31003/USPNF_M29350_05_01

DOI ref: x6eql