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Promethazine Hydrochloride Injection

DEFINITION

Promethazine Hydrochloride Injection is a sterile solution of Promethazine Hydrochloride in Water for Injection. It contains NLT 95.0% and NMT 110.0% of the labeled amount of promethazine hydrochloride ($C_{17}H_{20}N_2S \cdot HCl$).

[NOTE—Throughout the following procedures, protect the samples, the Reference Standards, and the solutions containing them, by conducting the procedures without delay under subdued light or using low-actinic glassware.]

IDENTIFICATION

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

ASSAY

• PROCEDURE

Buffer: 3.7 g/L of ammonium acetate in water

Solution A: Acetonitrile and *Buffer* (30:70)

Mobile phase: See [Table 1](#).

Table 1

Time (min)	Solution A (%)	Acetonitrile (%)
0	100	0
10	60	40
18	60	40
18.1	100	0
25	100	0

Diluent: 0.1% triethylamine in methanol

System suitability solution: 1.0 µg/mL each of [USP Promethazine Hydrochloride RS](#) and [USP Promethazine Related Compound B RS](#) in *Diluent*

Standard solution: 0.05 mg/mL of [USP Promethazine Hydrochloride RS](#) in *Diluent*

Sample solution: Nominally 0.05 mg/mL of promethazine hydrochloride from a volume of injection in *Diluent*

[NOTE—Sonication may be used in the preparation of *System suitability solution*, *Standard solution*, and *Sample solution*.]

Chromatographic system

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

Mode: LC

Detector: UV 254 nm. For *Identification B*, use a diode array detector in the range of 200–400 nm.

Column: 4.6-mm × 15-cm; 5-µm packing L1

Temperatures

Column: 30°

Autosampler: 4°

Flow rate: 1.4 mL/min

Injection volume: 15 µL

System suitability**Samples:** System suitability solution and Standard solution

[NOTE—The relative retention times for promethazine and promethazine related compound B are 1.0 and 1.3, respectively.]

Suitability requirements**Resolution:** NLT 5.0 between the promethazine and promethazine related compound B peaks, *System suitability solution***Tailing factor:** NMT 2.0, *Standard solution***Relative standard deviation:** NMT 1.0%, *Standard solution***Analysis****Samples:** Standard solution and Sample solutionCalculate the percentage of the labeled amount of promethazine hydrochloride ($C_{17}H_{20}N_2S \cdot HCl$) in the portion of Injection taken:

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

 r_U = peak response from the *Sample solution* r_S = peak response from the *Standard solution* C_S = concentration of [USP Promethazine Hydrochloride RS](#) in the *Standard solution* (mg/mL) C_U = nominal concentration of promethazine hydrochloride in the *Sample solution* (mg/mL)**Acceptance criteria:** 95.0%–110.0%**IMPURITIES****• ORGANIC IMPURITIES****Mobile phase, Diluent, and Chromatographic system:** Proceed as directed in the Assay.**Standard solution:** 1.0 μ g/mL each of [USP Promethazine Hydrochloride RS](#) and [USP Promethazine Related Compound B RS](#) in *Diluent***Sample solution:** Nominally 500 μ g/mL of promethazine hydrochloride from a volume of Injection in *Diluent*[NOTE—Sonication may be used in the preparation of *Standard solution* and *Sample solution*.]**System suitability****Sample:** Standard solution[NOTE—See [Table 2](#) for the relative retention times.]**Suitability requirements****Resolution:** NLT 5.0 between the promethazine and promethazine related compound B peaks**Relative standard deviation:** NMT 2.0% for promethazine**Analysis****Samples:** Standard solution and Sample solution

Calculate the percentage of each degradation product in the portion of Injection taken:

$$\text{Result} = (r_U/r_S) \times (C_S/C_U) \times (1/F) \times 100$$

 r_U = peak response of each degradation product from the *Sample solution* r_S = peak response of promethazine from the *Standard solution* C_S = concentration of [USP Promethazine Hydrochloride RS](#) in the *Standard solution* (μ g/mL) C_U = nominal concentration of promethazine hydrochloride in the *Sample solution* (μ g/mL) F = relative response factor (see [Table 2](#))**Acceptance criteria:** See [Table 2](#). Disregard peaks less than 0.05%.**Table 2**

Name	Relative Retention Time	Relative Response Factor	Acceptance Criteria, NMT (%)
Promethazine sulfoxide ^a	0.3	0.29	2.6
Desmethyl promethazine ^b	0.7	1.1	0.2
Promethazine	1.0	—	—
Promethazine related compound B ^c	1.3	—	—
Phenothiazine ^d	1.7	2.3	0.2
Any individual unspecified degradation product	—	1.0	0.1
Total degradation products	—	—	2.8

^a *N,N*-Dimethyl-1-(10*H*-phenothiazin-10-yl)propan-2-amine sulfoxide.

^b *N*-Methyl-1-(10*H*-phenothiazin-10-yl)propan-2-amine.

^c This is a process impurity that is controlled in the drug substance and is included for identification only.

^d 10*H*-Phenothiazine.

SPECIFIC TESTS

- **BACTERIAL ENDOTOXINS TEST (85):** NMT 5.0 USP Endotoxin Units/mg of promethazine hydrochloride
- **pH (791):** 4.0–5.5
- **OTHER REQUIREMENTS:** It meets the requirements in *Injections and Implanted Drug Products (1)*.

ADDITIONAL REQUIREMENTS

- **PACKAGING AND STORAGE:** Preserve in single-dose or in multiple-dose containers, preferably of Type I glass, protected from light.

- **USP REFERENCE STANDARDS (11).**

[USP Promethazine Hydrochloride RS](#)

[USP Promethazine Related Compound B RS](#)

Isopromethazine hydrochloride;

N,N-Dimethyl-2-(10*H*-phenothiazin-10-yl)propan-1-amine hydrochloride.

$C_{17}H_{20}N_2S \cdot HCl$

320.88

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

Topic/Question	Contact	Expert Committee
PROMETHAZINE HYDROCHLORIDE INJECTION	Documentary Standards Support	SM52020 Small Molecules 5
REFERENCE STANDARD SUPPORT	RS Technical Services RSTECH@usp.org	SM52020 Small Molecules 5

Chromatographic Database Information: [Chromatographic Database](#)

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