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# Meperidine Hydrochloride Tablets

## DEFINITION

Meperidine Hydrochloride Tablets contain NLT 95.0% and NMT 105.0% of the labeled amount of meperidine hydrochloride ( $C_{15}H_{21}NO_2 \cdot HCl$ ).

## IDENTIFICATION

### • A. ~~IDENTIFICATION—ORGANIC NITROGENOUS BASES (181)~~

**Sample solution:** Transfer an amount nominally equivalent to about 50 mg of meperidine hydrochloride from powdered Tablets to a separator, add 10 mL of water, and shake. Add 5 mL of saturated sodium chloride solution and 1 mL of sodium hydroxide solution (1 in 25). Extract with three 20-mL portions of chloroform, filtering the extracts through cotton overlaid with anhydrous sodium sulfate. Evaporate the chloroform on a steam bath, and dissolve the residue in 4 mL of carbon disulfide.

**Standard solution:** In a second separator, proceed as directed in the *Sample solution*, using 50 mg of [USP Meperidine Hydrochloride RS](#).

**Analysis:** Proceed as directed in the chapter, beginning with “Determine the absorption spectra”.

**Acceptance criteria:** Meet the requirements

### • 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

**Solution A:** Transfer about 6.8 g of monobasic potassium phosphate to a 1000-mL volumetric flask. Dissolve in and dilute with water to volume. Add 10 mL of triethylamine, and mix. Adjust with phosphoric acid to a pH of 7.0, and filter.

**Mobile phase:** Acetonitrile and *Solution A* (550:450), filtered and degassed

**Standard stock solution:** 0.6 mg/mL of [USP Meperidine Hydrochloride RS](#) in water

**Standard solution:** 0.12 mg/mL of [USP Meperidine Hydrochloride RS](#) from the *Standard stock solution* in *Mobile phase*

**Sample stock solution:** Transfer an amount nominally equivalent to about 60 mg of meperidine hydrochloride from NLT 20 finely powdered Tablets to a 100-mL volumetric flask. Add about 70 mL of *Mobile phase*, and sonicate for 10 min with occasional shaking. Shake by mechanical means for about 30 min, dilute with *Mobile phase* to volume, mix, and filter.

**Sample solution:** Nominally equivalent to 0.12 mg/mL of meperidine hydrochloride in *Mobile phase* from the *Sample stock solution*

### Chromatographic system

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

**Mode:** LC

**Detector:** UV 230 nm

**Column:** 3.9-mm × 30-cm; packing L1

**Flow rate:** 1 mL/min

**Injection volume:** 20 µL

### System suitability

**Sample:** *Standard solution*

### Suitability requirements

**Column efficiency:** NLT 2000 theoretical plates

**Tailing factor:** NMT 2 for the meperidine peak

**Relative standard deviation:** NMT 2%

### Analysis

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

Calculate the percentage of meperidine hydrochloride ( $C_{15}H_{21}NO_2 \cdot HCl$ ) in the portion of Tablets taken:

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

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

$r_s$  = peak response of meperidine from the *Standard solution*

$C_s$  = concentration of [USP Meperidine Hydrochloride RS](#) in the *Standard solution* (mg/mL)

$C_u$  = nominal concentration of meperidine hydrochloride in the *Sample solution* (mg/mL)

**Acceptance criteria:** 95.0%–105.0%

#### PERFORMANCE TESTS

• [DISSOLUTION \(711\)](#)

**Medium:** Water; 500 mL

**Apparatus 1:** 100 rpm

**Time:** 45 min

**Standard solution:** A known concentration of [USP Meperidine Hydrochloride RS](#) in *Medium*

**Sample solution:** Filter portions of the solution under test, and suitably dilute with *Medium*, if necessary.

**Chromatographic system and System suitability:** Proceed as directed in the Assay.

**Analysis:** Determine the labeled amount of meperidine hydrochloride ( $C_{15}H_{21}NO_2 \cdot HCl$ ) dissolved by comparing the peak response of meperidine from the *Sample solution* with that from the *Standard solution*.

**Tolerances:** NLT 75% (Q) of the labeled amount of meperidine hydrochloride ( $C_{15}H_{21}NO_2 \cdot HCl$ ) is dissolved.

• [UNIFORMITY OF DOSAGE UNITS \(905\)](#): Meet the requirements

#### ADDITIONAL REQUIREMENTS

• **PACKAGING AND STORAGE:** Preserve in well-closed, light-resistant containers.

• [USP REFERENCE STANDARDS \(11\)](#)

[USP Meperidine Hydrochloride RS](#)

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

Topic/Question	Contact	Expert Committee
MEPERIDINE HYDROCHLORIDE TABLETS	<a href="#">Documentary Standards Support</a>	SM22020 Small Molecules 2
REFERENCE STANDARD SUPPORT	RS Technical Services <a href="mailto:RSTECH@usp.org">RSTECH@usp.org</a>	SM22020 Small Molecules 2

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

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