

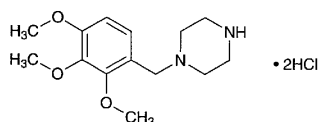
yphenidyl hydrochloride ($C_{20}H_{31}NO \cdot HCl$), dissolve in 2 mL of dilute hydrochloric acid and 60 mL of water by warming on a water bath for 10 minutes with occasional shaking. After cooling, add 2 mL of methanol and water to make exactly 100 mL, and use this solution as the sample solution. Dissolve about 0.05 g of Trihexyphenidyl Hydrochloride Reference Standard (determine previously its loss on drying at $105^\circ C$ for 3 hours), weighed accurately, in methanol, add methanol to make exactly 20 mL. Pipet 2 mL of this solution, add 2 mL of dilute hydrochloric acid and water to make exactly 100 mL, and use this solution as the standard solution. Pipet 10 mL each of the sample solution and the standard solution into glass-stoppered centrifuge tubes, add exactly 10 mL each of bromocresol purple-dipotassium hydrogenphosphate-citric acid TS and 15 mL each of chloroform, stopper tightly, shake thoroughly, and centrifuge. Pipet 10 mL each of the chloroform layers, and add chloroform to make exactly 50 mL. Determine the absorbances, A_T and A_S , of the subsequent solutions of the sample solution and the standard solution at 408 nm as directed under Spectrophotometry, respectively.

$$\begin{aligned} & \text{Amount (mg) of trihexyphenidyl hydrochloride} \\ & (C_{20}H_{31}NO \cdot HCl) \\ & = \text{amount (mg) of Trihexyphenidyl Hydrochloride} \\ & \quad \text{Reference Standard, calculated on the dried basis} \\ & \quad \times \frac{A_T}{A_S} \times \frac{1}{10} \end{aligned}$$

Containers and storage Containers—Tight containers.

Trimetazidine Hydrochloride

塩酸トリメタジジン



$C_{14}H_{22}N_2O_3 \cdot 2HCl$: 339.26
1-(2,3,4-Trimethoxybenzyl)piperazine dihydrochloride
[13171-25-0]

Trimetazidine Hydrochloride contains not less than 98.0% of $C_{14}H_{22}N_2O_3 \cdot 2HCl$, calculated on the dried basis.

Description Trimetazidine Hydrochloride occurs as a white, crystalline powder. It is odorless.

It is very soluble in water, sparingly soluble in ethanol (95), and practically insoluble in diethyl ether.

The pH of a solution of Trimetazidine Hydrochloride (1 in 20) is between 2.3 and 3.3.

Melting point: about $227^\circ C$ (with decomposition).

Identification (1) Dissolve 5 mg of Trimetazidine Hydrochloride in 1 mL of water, add 1 mL of *p*-benzoquinone TS, boil gently for 2 to 3 minutes, and cool: a red color develops.

(2) Determine the absorption spectrum of a solution of Trimetazidine Hydrochloride in 0.1 mol/L hydrochloric

acid TS (1 in 6250) 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.

(3) Determine the infrared absorption spectrum of Trimetazidine Hydrochloride as directed in the potassium chloride 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.

(4) A solution of Trimetazidine Hydrochloride (1 in 50) responds to the Qualitative Tests for chloride.

Purity (1) Clarity and color of solution—Dissolve 0.5 g of Trimetazidine Hydrochloride in 5 mL of water: the solution is clear and colorless.

(2) Heavy metals—Proceed with 2.0 g of Trimetazidine Hydrochloride according to Method 2, and perform the test. Prepare the control solution with 2.0 mL of Standard Lead Solution (not more than 10 ppm).

(3) Arsenic—Prepare the test solution with 1.0 g of Trimetazidine Hydrochloride according to Method 1, and perform the test using Apparatus B (not more than 2 ppm).

(4) Related substances—Dissolve 0.5 g of Trimetazidine Hydrochloride in 10 mL of water, and use this solution as the sample solution. Pipet 1 mL of the sample solution, add water to make exactly 100 mL, and use this solution as the standard solution. Perform the test with these solutions as directed under the Thin-layer Chromatography. Spot $10 \mu L$ 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 cyclohexane and diethylamine (1:1) to a distance of about 10 cm, air-dry the plate, and then dry at $110^\circ C$ for 1 hour. After cooling, examine under ultraviolet light (main wavelength: 254 nm): the spots other than the principal spot and the spot of the starting point from the sample solution are not more intense than the spot from the standard solution.

Water Not more than 1.5% (2 g, direct titration).

Residue on ignition Not more than 0.10% (1 g).

Assay Weigh accurately about 0.25 g of Trimetazidine Hydrochloride, dissolve in 20 mL of water, add 5 mL of sodium hydroxide TS, and extract with three 20-mL portions of chloroform. Filter the chloroform extract each time through a pledget of cotton with anhydrous sodium sulfate on a funnel. Combine all the extracts, and titrate with 0.1 mol/L perchloric acid VS until the color of the solution changes from red-brown to green (indicator: 0.5 mL of *p*-naphtholbenzein TS). Perform a blank determination, and make any necessary correction.

$$\begin{aligned} & \text{Each mL of 0.1 mol/L perchloric acid VS} \\ & = 16.963 \text{ mg of } C_{14}H_{22}N_2O_3 \cdot 2HCl \end{aligned}$$

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