Certificate of Analysis

Product Name: Terazosin hydrochloride

CAS Number: 63074-08-8 **IUPAC Name:**

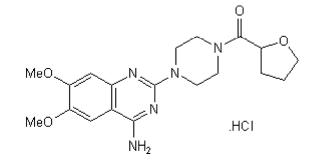
1-(4-Amino-6,7-dimethoxy-2-quinazolinyl)-4-[(tetrahydro-2-furanyl)carbonyl]-piperazine hydrochloride

1. PHYSICAL AND CHEMICAL PROPERTIES

Batch Molecular Formula: Batch Molecular Weight: Physical Appearance: Solubility:

Batch Molecular Structure:

C₁₉H₂₅N₅O₄.HCl.1³/₄H₂O 455.42 White solid water to 20 mM with gentle warming Store at RT



2. ANALYTICAL DATA

Storage:

HPLC:
¹ H NMR:
Mass Spectrum:
Microanalysis:

Shows 99.1% purity	
Consistent with structure	
Consistent with structure	

	Carbon Hy	/drogen N	litrogen	Chlorine
Theoretical	50.11	6.53	15.38	7.78
Found	49.49	6.63	15.13	7.54

Caution - Not Fully Tested • Research Use Only • Not For Human or Veterinary Use

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Print Date: Sep 27th 2024

Catalog No.: 1506 Batch No.: 2

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CAS Number: 63074-08-8

IUPAC Name: 1-(4-Amino-6,7-dimethoxy-2-quinazolinyl)-4-[(tetrahydro-2-furanyl)carbonyl]-piperazine hydrochloride

Description:

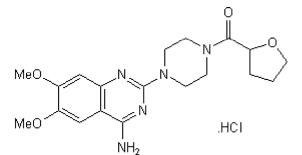
Terazosin hydrochloride is a selective α_1 antagonist. Increases phosphoglycerate kinase 1 (PGK1) activity (K_d = 2.9 μ M). Upregulates glucose metabolism and increases ATP production. In an in vitro model of GI disease, terazosin reduces inflammation and protects against oxidative damage. Neuroprotective in in vivo models of ALS and Parkinson's Disease. Antihypertensive following oral or intravenous administration in vivo.

Physical and Chemical Properties:

Batch Molecular Formula: $C_{19}H_{25}N_5O_4$.HCl.1³/₄H₂O Batch Molecular Weight: 455.42 Physical Appearance: White solid

Minimum Purity: ≥98%

Batch Molecular Structure:



Storage: Store at RT

Solubility & Usage Info:

water to 20 mM with gentle warming

Stability and Solubility Advice:

Some solutions can be difficult to obtain and can be encouraged by rapid stirring, sonication or gentle warming (in a 45-60°C water bath).

Catalog No.: 1506

Information concerning product stability, particularly in solution, has rarely been reported and in most cases we can only offer a general guide. *Unless contradicted by product-specific protocols or instructions, our standard recommendations apply:

SOLIDS: Provided storage is as stated on the product label and the vial is kept tightly sealed, the product can be stored for up to 6 months from date of receipt.

SOLUTIONS: We recommend that stock solutions, once prepared, are stored aliquoted in tightly sealed vials at -20°C or below and used within 1 month. Wherever possible solutions should be made up and used on the same day.

References:

Hancock *et al* (1995) Actions of tera. and its enantiomers at subtypes of α 1- and α 2-adrenoceptors *in vitro*. J.Recept.Signal Transduct.Res. **15** 863. PMID: 8673721.

Maruyama *et al* (1994) Comparison of displacemental potencies of tera. enantiomers for α_1 -adrenoceptor subtypes. Biol.Pharm.Bull. **17** 1126. PMID: 7820122.

Kyncl (1986) Pharmacology of tera. Am.J.Med. 80 12. PMID: 2872801.

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