

Certificate of Analysis

Print Date: Jan 13th 2016

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Product Name: Amthamine dihydrobromide Catalog No.: 0668 Batch No.: 2

CAS Number: 142457-00-9

IUPAC Name: 2-Amino-5-(2-aminoethyl)-4-methylthiazole dihydrobromide

1. PHYSICAL AND CHEMICAL PROPERTIES

Batch Molecular Formula: C₆H₁₁N₃S.2HBr

Batch Molecular Weight: 319.06 **Physical Appearance:** White solid

Solubility: water to 100 mM
Storage: Desiccate at -20°C

Batch Molecular Structure:

2. ANALYTICAL DATA

TLC: $R_f = 0.25$ (Dichloromethane:Methanol:Ammonia soln. [9:1:0.1])

Melting Point: At 275°C(dec)

HPLC: Shows >99.1% purity
 ¹H NMR: Consistent with structure
 Mass Spectrum: Consistent with structure

Microanalysis: Carbon Hydrogen Nitrogen

Theoretical 22.59 4.11 13.17 Found 22.8 4.06 12.82



Product Information

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IUPAC Name: 2-Amino-5-(2-aminoethyl)-4-methylthiazole dihydrobromide

Description:

Highly selective H_2 agonist, slightly more potent than histamine itself. Only a weak antagonist at H_3 and has no activity at H_1 receptors. Induces vasodilation of cerebral arteries and decreases myogenic tone in vitro.

Physical and Chemical Properties:

Batch Molecular Formula: $C_6H_{11}N_3S.2HBr$

Batch Molecular Weight: 319.06 Physical Appearance: White solid

Minimum Purity: >99%

Batch Molecular Structure:

Storage: Desiccate at -20°C

Solubility & Usage Info:

water to 100 mM

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).

Information concerning product stability, particularly in solution, has rarely been reported and in most cases we can only offer a general guide. Our standard recommendations are:

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:

Eriks et al (1992) Histamine H_2 -receptor agonists. Synthesis, in vitro pharmacology, and qualitative structure-activity relationships of substituted 4- and 5-(2-aminoethyl)thiazole. J.Med.Chem. **35** 3239. PMID: 1507209.

Coruzzi *et al* (1993) The new potent and selective histamine H₂ receptor agonist amthamine as a tool to study gastric secretion. Naunyn Schmiedebergs Arch.Pharmacol. *348* 77. PMID: 8377843.

Poli et al (1993) In vitro cardiac pharmacology of the new histamine H₂ receptor agonist amthamine; comparisons with histamine and dimaprit. Agents Actions **40** 44. PMID: 8147269.

Jarajapu *et al* (2006) Histamine decreases myogenic tone in rat cerebral arteries by H₂-receptor-mediated Kv channel activation, independent of endothelium and cyclic AMP. Eur.J.Pharmacol. *547* 116. PMID: 16920098.