

Certificate of Analysis

Print Date: Jan 15th 2016

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Product Name: MK 217 Catalog No.: 4014 Batch No.: 1

CAS Number: 129318-43-0

IUPAC Name: P,P-(4-Amino-1-hydroxybutylidene) bis-phosphonic acid monosodium salt

1. PHYSICAL AND CHEMICAL PROPERTIES

Batch Molecular Formula: $C_4H_{12}NNaO_7P_2.21/2H_2O$

Batch Molecular Weight: 316.12

Physical Appearance: White solid

Solubility: water to 50 mM

Storage: Desiccate at -20°C

Batch Molecular Structure:

2. ANALYTICAL DATA

¹H NMR: Consistent with structure Mass Spectrum: Consistent with structure

Microanalysis: Carbon Hydrogen Nitrogen

Theoretical 15.2 5.42 4.43 Found 15.02 5.55 4.34



Product Information

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IUPAC Name: P,P'-(4-Amino-1-hydroxybutylidene)bis-phosphonic acid monosodium salt

Description:

Osteoclast-mediated bone resorption inhibitor. Binds and blocks farnesyl diphosphate synthase (FPPS) in the HMG-CoA pathway ($IC_{50} = 460$ nM for recombinant human FPPS); causes macrophage apoptosis. Inhibits prenylation and sterol biosynthesis in purified osteoclasts.

Physical and Chemical Properties:

Batch Molecular Formula: C₄H₁₂NNaO₇P₂.2½H₂O

Batch Molecular Weight: 316.12 Physical Appearance: White solid

Batch Molecular Structure:

Storage: Desiccate at -20°C

Solubility & Usage Info:

water to 50 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:

Fisher et al (1999) Alendronate mechanism of action; geranylgeraniol, an intermediate in the mevalonate pathway, prevents inhibition of osteoclast formation, bone resorption, and kinase activation in vitro. Proc.Natl.Acad.Sci. 96 133.

Bergstrom et al (2000) Alendronate is a specific, nanomolar inhibitor of fanesyl diphosphate synthase. Arch.Biochem.Biophys. 373 231. PMID: 10620343.

Fournier et al (2008) Lowering bone mineral affinity of bisphosphonates as a therapeutic strategy to optimize skeletal tumor growth inhibition in vitro. Cancer Res. 68 8945. PMID: 18974139.