

Certificate of Analysis

Print Date: Jan 15th 2016

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Product Name: AG 99 Catalog No.: 0497 Batch No.: 1

CAS Number: 122520-85-8

IUPAC Name: (E)-2-Cyano-3-(3,4-dihydroxyphenyl)-2-propenamide

1. PHYSICAL AND CHEMICAL PROPERTIES

 $\begin{array}{lll} \textbf{Batch Molecular Formula:} & \textbf{C_{10}H}_{8}\textbf{N_{2}O}_{3}\\ \textbf{Batch Molecular Weight:} & 204.18\\ \textbf{Physical Appearance:} & \textbf{Yellow solid}\\ \textbf{Solubility:} & \textbf{ethanol to 5 mM} \end{array}$

DMSO to 100 mM

Storage: Desiccate at +4°C

Batch Molecular Structure:

HO CN NH

2. ANALYTICAL DATA

TLC: $R_f = 0.65$ (Dichloromethane:Methanol [10:1])

HPLC: Shows >99.4% purity

¹H NMR: Consistent with structure

Mass Spectrum: Consistent with structure

Microanalysis: Carbon Hydrogen Nitrogen

Theoretical 58.82 3.95 13.71 Found 58.63 3.91 13.62



Product Information

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IUPAC Name: (E)-2-Cyano-3-(3,4-dihydroxyphenyl)-2-propenamide

Description:

Epidermal growth factor receptor (EGFR) kinase inhibitor (IC $_{\rm 50}$ = 10 $\,\mu$ M) that is selective over insulin receptor kinase. Blocks tyrosine phosphorylation of p145 $^{\rm met}$ and promotes cell death of normal and cancer cells via activation of caspase-like proteases in vitro.

Physical and Chemical Properties:

Batch Molecular Formula: C₁₀H₈N₂O₃ Batch Molecular Weight: 204.18 Physical Appearance: Yellow solid

Minimum Purity: >99%

Batch Molecular Structure:

Storage: Desiccate at +4°C

Solubility & Usage Info:

ethanol to 5 mM DMSO 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:

Gazit et al (1989) Tyrphostins. I. Synthesis and biological activity of protein tyrosine kinase inhibitors. J.Med.Chem. 32 2344. PMID: 2552117.

Levitzki and Gilon (1991) Tyrphostins as molecular tools and potential antiproliferative drugs. TiPS 12 171. PMID: 1862529.

Gazit (1996) Tyrphostins. 6. Dimeric benzylidenemalononitrile tyrphostins: potent inhibitors of EGF receptor tyrosine kinase *in vitro*. J.Med.Chem. **39** 4905. PMID: 8960549.

Yamamoto *et al* (2006) Tyrosine phosphorylation of p145^{met} mediated by EGFR and Src is required for serum-independent survival of human bladder carcinoma cells. J.Cell Sci. *119* 4623. PMID: 17062641.