PRODUCT INFORMATION



I-OMe AG-538 Item No. 35090

CAS Registry No.: 1094048-77-7

Formal Name: 3,4-dihydroxy-aE-[(4-hydroxy-3-iodo-

5-methoxyphenyl)methylene]-β-oxo-

benzenepropanenitrile

Synonym: I-OMe Tyrphostin AG-538

MF: $C_{17}H_{12}INO_5$ FW: 437.2 **Purity:** ≥95% Supplied as: A solid Storage: -20°C Stability: ≥4 years

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

Laboratory Procedures

I-OMe AG-538 is supplied as a solid. A stock solution may be made by dissolving the I-OMe AG-538 in the solvent of choice, which should be purged with an inert gas. I-OMe AG-538 is soluble (≥10 mg/ml) in DMSO.

Description

I-OMe AG-538 is an inhibitor of the insulin-like growth factor 1 receptor (IGF-1R; IC $_{50}$ = 3.4 μ M) and phosphatidylinositol 5-phosphate 4-kinase α (PI5P4K α ; IC $_{50}$ = 1 μ M). 1,2 It is selectively cytotoxic to nutrient-deprived PANC-1 pancreatic cancer cells over nutrient-sufficient PANC-1 cells.³

References

- 1. Blum, G., Gazit, A., and Levitzki, A. Substrate competitive inhibitors of IGF-1 receptor kinase. Biochemistry **39(51)**, 15705-15712 (2000).
- 2. Davis, M.I., Sasaki, A.T., Shen, M., et al. A homogeneous, high-throughput assay for phosphatidylinositol 5-phosphate 4-kinase with a novel, rapid substrate preparation. PLoS One 8(1), e54127 (2013).
- 3. Momose, I., Kunimoto, S., Osono, M., et al. Inhibitors of insulin-like growth factor-1 receptor tyrosine kinase are preferentially cytotoxic to nutrient-deprived pancreatic cancer cells. Biochem. Biophys. Res. Commun. 380(1), 171-176 (2009).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

WARRANTY AND LIMITATION OF REMEDY

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