
Product Data Sheet

Product Name: Rac GTPase fragment
 Cat. No.: GP10032

Chemical Properties

Cas. No.

SMILES NC(C(C)C)C(NC(CC1=CC=CC=C1)C(NC(CC(O)=O)C(NC(CCC(O)=O)C(NC(C)C(NC(C(NC(CCCNC(N)=N)C(NC(C)C(NC(C(C)C)C(O)=O)=C

Formula C₄₆H₇₄N₁₂O₁₄

M.Wt

1019.15

Solubility ≥ 101.9mg/mL in DMSO

Storage

Store at -20°C

General tips For obtaining a higher solubility , please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be months.

Shipping Condition Evaluation sample solution : ship with blue ice All other available size: ship with RT , or blue ice upon request.

Structure

Background

Rac GTPase fragment, (C₄₆H₇₄N₁₂O₁₄), a peptide with the sequence H₂N-Val-Phe-Asp-Glu-Ala-Ile-Arg-Ala-Val-OH, MW= 1019.15. Rac is a subfamily of the Rho family of GTPases, small signaling G proteins (more specifically a GTPase). Rac1 is thought to play a significant role in the development of various cancers, including melanoma¹ and non-small cell lung cancer². Rac1 is a small (~21 kDa) signaling G protein (more specifically a GTPase), and is a member of the Rac subfamily of the family Rho family of GTPases. Members of this superfamily appear to regulate a diverse array of cellular events, including the control of cell growth, cytoskeletal reorganization, and the activation of protein kinases Rac1 is a pleiotropic regulator of many cellular processes, including the cell cycle, cell-cell adhesion, motility (through the actin network), and of epithelial differentiation³.

References:

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- Stallings-Mann, M. L.; Waldmann, J.; Zhang, Y.; Miller, E.; Gauthier, M. L.; Visscher, D. W.; Downey, G. P.; Radisky, E. S. et al. (2012). "Matrix Metalloproteinase Induction of Rac1b, a Key Effector of Lung Cancer Progression". Science Translational Medicine 4 (142): 142ra95-142ra95.
- Ridley A. (2006). "Rho GTPases and actin dynamics in membrane protrusions and vesicle trafficking". Trends Cell Biol 16 (10): 522-9.

Caution: Product has not been fully validated for medical applications. For research use only.

Tel: (909) 407-4943 Fax: (626) 353-8530 E-mail: tech@glpbio.com

Address: 10292 Central Ave. #205, Montclair, CA, USA