PRODUCT INFORMATION



• 10 NH₄+

D-myo-Inositol-1,3,4,5,6-pentaphosphate (ammonium salt)

Item No. 10009851

Formal Name: D-myo-inositol-1,3,4,5,6-penta

(hydrogen phosphate), decaammonium salt

Ins(1,3,4,5,6)P₅, Synonyms:

1,3,4,5,6-IP₅ (sodium salt)

MF: $C_6H_7O_{21}P_5 \bullet 10NH_4$

FW: 750.4 **Purity:** ≥98%

Supplied as:

Storage: -20°C Stability:

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

A lyophilized powder ≥5 years

Laboratory Procedures

D-myo-Inositol-1,3,4,5,6-pentaphosphate (Ins(1,3,4,5,6) P_s) (ammonium salt) is supplied as a lyophilized powder. A stock solution may be made by dissolving the Ins(1,3,4,5,6)P₅ (ammonium salt) in water. The solubility of Ins(1,3,4,5,6)P₅ (ammonium salt) in water is approximately 50 mg/ml. We do not recommend storing the aqueous solution for more than one day.

Description

 $lns(1,3,4,5,6)P_5$ is one of the many inositol phosphate isomers that act as small, soluble second messengers in the transmission of cellular signals.¹⁻³ It can be interconverted with Ins(3,4,5,6)P₄ by a 1-kinase/1-phosphatase cycle, as well as with Ins(1,4,5,6)P₄ in a 3-kinase/3-phosphatase cycle.³ Ins(1,3,4,5,6)P₅ inhibits the phosphorylation and kinase activity of Akt/PKB, inducing apoptosis in ovarian, lung, and breast cancer cells.⁴ It exhibits antiangiogenic activity in vitro, blocking capillary tube formation of HUVEC, as well as antitumor effects against cancer xenografts in nude mice. Ins(1,3,4,5,6)P₅ binds to the PH domain of Grp1 with a K_d of 590 nM.⁶

References

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- Shears, S.B. The versatility of inositol phosphates as cellular signals. Biochim. Biophys. Acta. 1436(1-2), 49-67 (1998).
- 4. Piccolo, E., Vignati, S., Maffucci, T., et al. Inositol pentakisphosphate promotes apoptosis through the PI 3-K/Akt pathway. Oncogene 23(9), 1754-1765 (2004).
- 5. Maffucci, T., Piccolo, E., Cumashi, A., et al. Inhibition of the phosphatidylinositol 3-kinase/Akt pathway by inositol pentakisphosphate results in antiangiogenic and antitumor effects. Cancer Res. **65(18)**, 8339-8349 (2005).
- 6. Kavran, J.M., Klein, D.E., Lee, A., et al. Specificity and promiscuity in phosphoinositide binding by pleckstrin homology domains. J. Biol. Chem. 273(46), 30497-30508 (1998).

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.

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