

Product Information

Phospholipase A₂ from honey bee venom (*Apis mellifera*)

Catalog Number **P9279**
Storage Temperature -20 °C

CAS RN 9001-84-7

EC 3.1.1.4

Synonyms: Phosphatidylcholine 2-acylhydrolase,
Lecithinase A, PLA2

Product Description

Molecular mass:¹ 14.5 kDa (amino acid sequence)
pI:² 10.5 ± 1.0

Phospholipase A₂ reacts stereospecifically with most *sn*-3-phosphoglycerides. The fatty acid ester bonds are hydrolyzed at the C-2 position. This reaction requires calcium for catalysis. The general reaction catalyzed is:

phosphatidylcholine + H₂O →
1-acylglycerophosphocholine + fatty acid

Phospholipase A₂ is inhibited *in vitro* by both calpactin I and calpactin II. The calpactins sequester the phospholipid substrate. There is no direct interaction between the calpactins and phospholipase A₂.³

Quinacrine has also been described as an inhibitor of phospholipase A₂ (IC₅₀ = 17 μM).⁴

Phospholipase A₂ is a single polypeptide chain of ~123 amino acids containing seven disulfide bridges. The crystal structure of phospholipase A₂ from bee venom, complexed with a phosphonate transition-state analogue compound, has been reported.⁵

Precautions and Disclaimer

This product is for R&D use only, not for drug, household, or other uses. Please consult the Safety Data Sheet for information regarding hazards and safe handling practices.

Preparation Instructions

Sigma-Aldrich tests the solubility of this product at 1.0 mg/mL in water. One publication reports the preparation of stock solutions of this product at the equivalent of 17.5 mg/mL in an aqueous buffer that contains 10 mM MOPS, pH 7.2, 0.5 mM EGTA, 100 mM HCl, and 4 mM CaCl₂.⁶

References

1. De Haas, G.H. *et al.*, Studies on phospholipase A and its zymogen from porcine pancreas. II. The assignment of the position of the six disulfide bridges. *Biochim. Biophys. Acta*, **221(1)**, 54-61 (1970).
2. De Haas, G.H. *et al.*, Purification and properties of phospholipase A from porcine pancreas. *Biochim. Biophys. Acta*, **159(1)**, 103-117 (1968).
3. Davidson, F.F. *et al.*, Inhibition of phospholipase A2 by "lipocortins" and calpactins. An effect of binding to substrate phospholipids. *J. Biol. Chem.*, **262(4)**, 1698-1705 (1987).
4. Magolda, R.L. *et al.*, "Novel Synthesis of Potent Site-Specific Phospholipase A2 Inhibitors", from *Prostaglandins, Leukotrienes and Lipoxins* (J.M. Bailey, ed). Plenum Press (New York, NY), pp. 669-672 (1985).
5. Scott, D.L. *et al.*, Crystal structure of bee-venom phospholipase A2 in a complex with a transition-state analogue. *Science*, **250(4987)**, 1563-1566 (1990).
6. Bollinger, J.G. *et al.*, Improved method for the quantification of lysophospholipids including enol ether species by liquid chromatography-tandem mass spectrometry. *J. Lipid Res.*, **51(2)**, 440-447 (2010).

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