

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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PRODUCT INFORMATION



(S)-Bromoenol lactone-d₇

Item No. 10535

Formal Name: 6E-(bromoethylene)tetrahydro-3S-

(1-naphthalenyl-2,3,4,5,6,7,8-d₇)-2H-

pyran-2-one

Synonym: (S)-BEL-d₇ MF: C₁₆H₆BrD₇O₂

FW:

Chemical Purity: ≥98% ((S)-Bromoenol lactone)

Deuterium

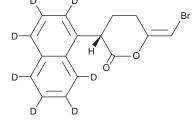
Incorporation: ≥99% deuterated forms (d_1-d_7) ; ≤1% d_0

UV/Vis.: λ_{max} : 280 nm

A solution in methyl acetate Supplied as:

-20°C Storage: ≥2 years Stability:

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



Laboratory Procedures

(S)-Bromoenol lactone-d₇ ((S)-BEL-d₇) is intended for use as an internal standard for the quantification of (S)-BEL by GC- or LC-MS. The accuracy of the sample weight in this vial is between 5% over and 2% under the amount shown on the vial. If better precision is required, the deuterated standard should be quantitated against a more precisely weighed unlabeled standard by constructing a standard curve of peak intensity ratios (deuterated versus unlabeled).

(S)-BEL-d₇ is supplied as a solution in methyl acetate. To change the solvent, simply evaporate the methyl acetate under a gentle stream of nitrogen and immediately add the solvent of choice. Solvents such as ethanol, DMSO, and dimethyl formamide purged with an inert gas can be used. The solubility of (S)-BEL-d₇ in these solvents is approximately 5, 25, and 50 mg/ml, respectively.

Description

The phospholipases are an extensive family of lipid hydrolases that function in cell signaling, digestion, membrane remodeling, and as venom components. The calcium-independent phospholipase A_2 (iPLA₂) are a PLA₂ subfamily closely associated with the release of arachidonic acid in response to physiologic stimuli. (S)-BEL) is an irreversible, chiral, mechanism-based inhibitor of iPLA $_{\beta}$ B that inhibits the vasopressin-induced release of arachidonate from cultured rat aortic smooth muscle (A10) cells with an IC_{50} value of 2 μ M.² (S)-BEL is more than 1,000-fold selective for iPLA₂ versus cPLA₂, and is 10-fold selective for iPLA₂β versus iPLA₂γ.

References

- 1. Balsinde, J., Balboa, M.A., Insel, P.A., et al. Regulation and inhibition of phospholipase A₂. Annu. Rev. Pharmacol. Toxicol. 39, 175-189 (1999).
- 2. Jenkins, C.M., Han, X., Mancuso, D.J., et al. Identification of calcium-independent phospholipase A2 (iPLA₂) β , and not iPLA₂ γ , as the mediator of arginine vasopressin-induced arachidonic acid release in A-10 smooth muscle cells. J. Biol. Chem. 277(36), 32807-32814 (2002).

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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