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Product Information



Bromoenol lactone-d-

Item No. 9000528

Formal Name: 6E-(bromomethylene)tetrahydro-3-(1-

naphthalenyl-2,3,4,5,6,7,8-d₇)-2H-pyran-2-one

Synonyms: BEL-d₇, Haloenol lactone-d₇, HELSS-d₇

MF: $C_{16}H_6D_7BrO_2$

FW: 324.2 **Chemical Purity:** ≥98%

Deuterium

 \geq 99% deuterated forms (d₁-d₇); \leq 1% d₀ Incorporation:

Stability: ≥1 year at -20°C

Supplied as: A solution in methyl acetate

UV/Vis.: λ_{max} : 223, 280 nm

Laboratory Procedures

Bromoenol lactone-d₇ (BEL-d₇) contains seven deuterium atoms at the 2, 3, 4, 5, 6, 7, and 8 positions. It is intended for use as an internal standard for the quantification of BEL by GC- or LC-mass spectrometry (MS). For long term storage, we suggest that BEL-d₇ be stored as supplied at -20°C. It should be stable for at least one year.

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 BEL-d₇ in these solvents is approximately 5, 25, and 50 mg/ml, respectively.

BEL-d₇ is used as an internal standard for the quantification of BEL by stable isotope dilution 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).

BEL is a selective, potent, irreversible, mechanism-based inhibitor of myocardial cytosolic calcium-independent phospholipase A₂ (iPLA₂) with a K₂ value of 180 nM. It also inhibits macrophage iPLA₂ in a concentration-dependent manner with an IC50 value of 60 nM and is an effective enzyme-activated irreversible inhibitor of chymotrypsin $(K_1 = 636 \text{ nM}).^{2,3}$

References

- 1. Hazen, S.L., Zupan, L.A., Weiss, R.H., et al. Suicide inhibition of canine myocardial cytosolic calcium-independent phospholipase A₂. J. Biol. Chem. 266, 7227-7232 (1991).
- 2. Ackermann, E.J., Conde-Frieboes, K., and Dennis, E.A. Inhibition of macrophage Ca²⁺-independent phospholipase A₂ by bromoenol lactone and trifluoromethyl ketones. J. Biol. Chem. 270, 445-450 (1995).
- Daniels, S.B., Cooney, E., Sofia, M.J., et al. Haloenol Lactones. Potent enzyme-activated irreversible inhibitors for α-chymotrypsin. J. Biol. Chem. 258, 15046-15053 (1983).

Related Products

For a list of related products please visit: www.caymanchem.com/catalog/9000528

WARNING: This product is for laboratory research only: not for administration to humans. Not for human or veterinary DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until information to the contrary becomes available. Do not ingest, swallow, or inhale. Do not get in eyes, on skin, or on clothing. Wash thoroughly after handling. This information contains some, but not all, of the information required for the safe and proper use of this material. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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