

Produktinformation



Forschungsprodukte & Biochemikalien



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Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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Product Information



(±)18-HETE

Item No. 10010638

CAS Registry No.: 133268-58-3

Formal Name: (±)18-hydroxy-5Z,8Z,11Z,14Z-

eicosatetraenoic acid

MF: $C_{20}H_{32}O_3$ FW: 320.5 **Purity:** ≥97%

Stability: ≥1 year at -20°C Supplied as: A solution in ethanol

Laboratory Procedures

For long term storage, we suggest that (±)18-HETE be stored as supplied at -20°C. It should be stable for at least one

 $(\pm)18$ -HETE is supplied as a solution in ethanol. To change the solvent, simply evaporate the ethanol under a gentle stream of nitrogen and immediately add the solvent of choice. Solvents such as DMSO and dimethyl formamide purged with an inert gas can be used. (±)18-HETE is miscible in these solvents.

Further dilutions of the stock solution into aqueous buffers or isotonic saline should be made prior to performing biological experiments. Ensure that the residual amount of organic solvent is insignificant, since organic solvents may have physiological effects at low concentrations. If an organic solvent-free solution of (±)18-HETE is needed, it can be prepared by evaporating the ethanol and directly dissolving the neat oil in aqueous buffers. The solubility of (±)18-HETE in PBS, pH 7.2, is approximately 0.8 mg/ml. For greater aqueous solubility, (±)18-HETE can be directly dissolved in 0.1 M Na₂CO₃ (solubility of approximately 2 mg/ml) and then diluted with PBS (pH 7.2) to achieve the desired concentration or pH. We do not recommend storing the aqueous solution for more than one day.

(±)18-HETE is the racemic version of a cytochrome P450 (CYP450) metabolite of arachidonic acid. When formed by the CYP2E1 isoform, 18-HETE is comprised 100% of the (R) isomer. 18(R)-HETE dose-dependently stimulates vasodilation of the rabbit kidney, whereas 18(S)-HETE does not affect perfusion pressure.² 18-HETE has negligible effects on ATPase activity.² 18(R)-HETE at 1 μM completely blocks 20-HETE-induced vasoconstriction of renal arterioles.³

References

- 1. Laethem, R.M., Balazy, M., Falck, J.R., et al. Formation of 19(S)-, 19(R)-, and 18(R)-hydroxyeicosatetraenoic acids by alcohol-inducible cytochrome P450 2E1. J. Biol. Chem. 268(17), 12912-12918 (1993).
- Carroll, M.A., Balazy, M., Margiotta, P., et al. Cytochrome P-450-dependent HETEs: Profile of biological activity and stimulation by vasoactive peptides. Am. J. Physiol. 271, R863-R869 (1996).
- Zhang, F., Deng, H., Kemp, R., et al. Decreased levels of cytochrome P450 2E1-derived eicosanoids sensitize renal arteries to constrictor agonists in spontaneously hypertensive rats. Hypertension 45, 103-108 (2005).

(±)17,18-DiHETE - Item No. 10006999 • (±)16-HETE - Item No. 10010635 • (±)17-HETE - Item No. 10010636 • 18(R)-HETE - Item No. 10010639 • 18(S)-HETE - Item No. 10010640

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 Material Safety Data Sheet, which has been sent via email to your institution.

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