Product Information



(R)-Flurbiprofen

Item No. 70255

CAS Registry No.: 51543-40-9

Formal Name: (R)-(-)-2-fluoro- α -methyl-4-

biphenylacetic acid

Synonyms: E-7869, Flurizan, Tarenflurbil

MF: $C_{15}H_{13}FO_2$ FW: 244.3 **Purity:** ≥99%

Stability: ≥2 year at room temperature

Supplied as: A crystalline solid UV/Vis.: λ_{max} : 247 nm

Laboratory Procedures

For long term storage, we suggest that (R)-flurbiprofen be stored as supplied at room temperature. It should be stable for at least two years.

(R)-Flurbiprofen is supplied s a crystalline solid. A stock solution may be made by dissolving the (R)-flurbiprofen in an organic solvent purged with an inert gas. (R)-Flurbiprofen is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF). The solubility of (R)-flurbiprofen in these solvents is approximately 10 mg/ml in DMSO and 25 mg/ml in ethanol and DMF.

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. Organic solvent-free aqueous solutions of (R)-flurbiprofen can be prepared by directly dissolving the crystalline compound in aqueous buffers. The solubility of (R)-flurbiprofen in PBS, pH 7.2, is approximately 0.9 mg/ml. We do not recommend storing the aqueous solution for more than one day.

(R)-Flurbiprofen is a member of the 2-aryl propionic acid group of nonsteroidal anti-inflammatory drugs (NSAIDs). Only a small amount (<5%) of (R)-enantiomer is converted to the (S)-enantiomer in rats and humans; therefore, the biological effects are specific to each enantiomer. Although inactive as an inhibitor of cyclooxygenase (COX), this enantiomer reduces inflammation through inhibition of NF-κB and AP-1 activation.² (R)-Flurbiprofen has also been shown to suppress prostate tumor cells by inducing p75NTR protein expression.³ (R)-Flurbiprofen inhibits the enzyme γ -secretase thereby preventing the formation of the amyloid β peptide (A β 42) from amyloid β precursor protein (APP). Before being dropped as a drug candidate, (R)-flurbiprofen advanced to Phase III clinical trials, the first drug candidate to advance to late stage trials for the treatment of mild Alzheimer's disease.

References

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- Tegeder, I., Niederberger, E., Israr, E., et al. Inhibition of NF-KB and AP-1 activation by R- and S-flurbiprofen1,2. FASEB J. 15, 595-597 (2001).
- Quann, E.J., Khwaja, F., Zavitz, K.H., et al. The aryl propionic acid R-flurbiprofen selectively induces p75NTRdependent decreased survival of prostate tumor cells. Cancer Res 67(7), 3254-3262 (2007).
- Kukar, T.L., Ladd, T.B., Bann, M.A., et al. Substrate-targeting g-secretase modulators. Nature 453, 925-929 (2008).

Related Products

For a list of related products please visit: www.caymanchem.com/catalog/70255

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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Said refund or replacement is conditioned on Buyer giving written notice to Cayman within thirty (30) days after arrival of the material at its destination. Failure of Buyer to give said notice within thirty (30) days shall constitute a waiver by Buyer of all claims hereunder with respect to said material.

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Cayman Chemical

Mailing address

1180 E. Ellsworth Road Ann Arbor, MI 48108 USA

Phone

(800) 364-9897 (734) 971-3335

(734) 971-3640

custserv@caymanchem.com

www.caymanchem.com