PRODUCT INFORMATION



Prostaglandin F_{2a} ethyl amide

Item No. 16016

CAS Registry No.: 54130-36-8

Formal Name: N-ethyl-9a,11a,15S-trihydroxy-

prosta-5Z,13E-dien-1-amide

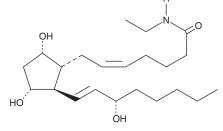
Synonyms: Dinoprost ethyl amide, PGF_{2a}-NEt

MF: $C_{22}H_{39}NO_4$ FW: 381.6 **Purity:** ≥98%

Supplied as: A solution in methyl acetate

Storage: -20°C Stability: ≥2 vears

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



Laboratory Procedures

Prostaglandin $F_{2\alpha}$ ethyl amide (PGF $_{2\alpha}$ -NEt) 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 PGF₂₀-NEt in these solvents is approximately 20, 16, and 33 mg/ml, respectively. PGF_{2n}-NEt is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, the methyl acetate solution of $PGF_{2\alpha}$ -NEt should be diluted with the aqueous buffer of choice. The solubility of PGF_{2a}-NEt in PBS (pH 7.2) is approximately 0.3 mg/ml. We do not recommend storing the aqueous solution for more than one day.

Description

 $PGF_{2\alpha}$ -NEt is an analog of $PGF_{2\alpha}$ in which the C-1 carboxyl group has been modified to an N-ethyl amide. PG esters have been shown to have ocular hypotensive activity. PG N-ethyl amides were recently introduced as alternative prostaglandin ocular hypotensive prodrugs.² Although it has been claimed that PG ethyl amides are not converted to the free acids in vivo, studies in our laboratories have shown that bovine and human corneal tissue converts the N-ethyl amides of various prostaglandins to the free acids with a conversion rate of about 2.5 μ g/g corneal tissue/hr.^{2,3} PGF_{2 α}-NEt would be expected to show the typical intraocular effects of PGF_{2 α} free acid, but with the much slower hydrolysis pharmacokinetics of the PG N-amides.

References

- 1. Bito, L.Z. Comparison of the ocular hypotensive efficacy of eicosanoids and related compounds. Exp. Eye Res. 38(2), 181-184 (1984).
- 2. Woodward, D.F., Krauss, A.H., Chen, J., et al. The pharmacology of bimatoprost (Lumigan™). Surv. Ophthalmol. 45(Suppl. 4), S337-S345 (2001).
- 3. Maxey, K.M., Johnson, J., and LaBrecque, J. The hydrolysis of bimatoprost in corneal tissue generates a potent prostanoid FP receptor agonist. Surv. Ophthalmol. 47(Suppl. 1), S34-S40 (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.

WARRANTY AND LIMITATION OF REMEDY

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