

# Produktinformation



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# **Product Information**



# 17-phenyl trinor Prostaglandin F $_{2\alpha}$ 1,15-lactone

Item No. 13992

Formal Name: 9α,11α,15S-trihydroxy-17-phenyl-

18,19,20-trinor-prosta-5Z,13E-dien-1-

oic acid, 1,15-lactone

Synonyms: Bimatoprost free acid 1,15 lactone,

17-phenyl trinor PGF $_{2\alpha}$  1,15-lactone

MF:  $C_{23}H_{30}O_4$ FW: 370.5 ≥98% **Purity:** 

Stability: ≥2 years at -20°C Supplied as: A crystalline solid

## **Laboratory Procedures**

For long term storage, we suggest that 17-phenyl trinor prostaglandin  $F_{2\alpha}$  1,15-lactone (17-phenyl trinor PGF<sub>2\alpha</sub> 1,15-lactone) be stored as supplied at -20°C. It should be stable for at least two years.

17-phenyl trinor  $PGF_{2\alpha}$  1,15-lactone is supplied as a crystalline solid. A stock solution may be made by dissolving the 17-phenyl trinor  $PGF_{2\alpha}$  1,15-lactone in the solvent of choice. 17-phenyl trinor  $PGF_{2\alpha}$  1,15-lactone is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide, which should be purged with an inert gas. The solubility of 17-phenyl trinor  $PGF_{2\alpha}$  1,15-lactone in these solvents is approximately 16, 12.5, and 14 mg/ml, respectively.

17-phenyl trinor PGF<sub>20</sub> 1,15-lactone is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, 17-phenyl trinor PGF<sub>20</sub>, 1,15-lactone should first be dissolved in ethanol and then diluted with the aqueous buffer of choice. 17-phenyl trinor  $PGF_{2\alpha}$  1,15-lactone has a solubility of approximately 0.5 mg/ml in a 1:1 solution of ethanol:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

17-phenyl trinor  $PGF_{2\alpha}$  (Item No. 16810) is a metabolically stable analog of  $PGF_{2\alpha}$  (Item No. 16010) with potent FP receptor agonist activity and well known intraocular pressure-reducing effects. 1-3 17-phenyl trinor  $PF_{2\alpha}$  1,15-lactone is the 1,15 lactone of 17-phenyl-trinor PGF<sub>20</sub>. Selective F-series PG derivatives such as this compound have been developed for suitable pharmacologic activity and an improved side-effect profile over current glaucoma therapeutics. 4.5

## References

- 1. Balapure, A.K., Rexroad, C.E., Jr., Kawada, K., et al. Structural requirements for prostaglandin analog interaction with the ovine corpus luteum prostaglandin  $F_{2\alpha}$  receptor. Biochem. Pharmacol. 38, 2375-2381 (1989).
- 2. Lake, S., Gullberg, H., Wahlqvist, J., et al. Cloning of the rat and human prostaglandin  $F_{2\alpha}$  receptors and the expression of the rat prostaglandin  $F_{2\alpha}$  receptor. FEBS Lett. 355, 317-325 (1994).
- Woodward, D.F., Krauss, A.H.-P., Chen, J., et al. The pharmacology of Bimatoprost (Lumigan™). Surv. Ophthalmol. 45, S337-S345 (2001).
- Stjernschantz, J.W. From  $PGF_{2\alpha}$ -isopropyl ester to latanoprost: A review of the development of xalatan. The proper lecture. Invest. Ophthamol. Vis. Sci. 42(6), 1134-1145 (2001).
- Maxey, K.M. and Stanton, M.L. Internal 1,15-lactones of fluprostenol and related prostaglandin  $F_{2\alpha}$  analogs and their use in the treatment of glaucoma and intraocular hypertension. US Patent 7,674,921 B2 (2010), 12/036,349.

## Related Products

For a list of related products please visit: www.caymanchem.com/catalog/13992

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

#### MATERIAL SAFETY DATA

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