

# Produktinformation



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Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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# PRODUCT INFORMATION



## 11β-Prostaglandin E<sub>2</sub>

Item No. 14510

CAS Registry No.: 38310-90-6

9-oxo-11β,15S-dihydroxy-prosta-5Z,13E-Formal Name:

dien-1-oic acid

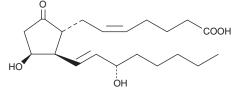
11β-PGE<sub>2</sub>, 11-epi PGE<sub>2</sub> Synonyms:

MF:  $C_{20}H_{32}O_5$ FW: 352.5 **Purity:** ≥98%

Supplied as: A solution in methyl acetate

Storage: -20°C Stability: ≥1 year

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



### **Laboratory Procedures**

11β-Prostaglandin E<sub>2</sub> (11β-PGE<sub>2</sub>) 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 11β-PGE<sub>2</sub> in these solvents is approximately 100 mg/ml.

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  $11\beta$ -PGE<sub>2</sub> is needed, it can be prepared by evaporating the methyl acetate and directly dissolving the neat oil in aqueous buffers. The solubility of  $11\beta$ -PGE<sub>2</sub> in PBS (pH 7.2) is approximately 5 mg/ml. We do not recommend storing the aqueous solution for more than one day.

#### Description

 $11\beta$ -PGE $_2$  is the C-11 epimer of PGE $_2$ . It is a moderate inhibitor of PGE $_2$  binding to rat hypothalamic membranes with a  $K_i$  value of 53 nM.  $^1$   $11\beta$ -PGE  $_2$  also stimulates bone resorption in rats at concentrations of 10 to 1,000 nM which is similar to  $PGE_2$ ,  $^2$  11 $\beta$ - $PGE_2$  inhibits  $PGE_2$  binding to the prostaglandin transporter protein with a K<sub>i</sub> of 56 nM.<sup>3</sup>

#### References

- 1. Dray, F. and Heaulme, M. Prostaglandins of the E series inhibit release of noradrenaline in rat hypothalamus by a mechanism unrelated to classical α<sub>2</sub> adrenergic presynaptic inhibition. Neuropharmacology 23(4), 457-462 (1984).
- 2. Raisz, L.G. and Woodiel, F.N. Effect of alterations in the cyclopentane ring on bone resorptive activity of prostaglandin. Prostaglandins 37(2), 229-235 (1989).
- Itoh, S., Lu, R., Bao, Y., et al. Structural determinants of substrates for the prostaglandin transporter PGT. Mol. Pharmacol. 50(4), 736-742 (1996).

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.

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