Product Information



Prostaglandin I₂ (sodium salt)

Item No. 18220

CAS Registry No.: 61849-14-7

 $6,9\alpha$ -epoxy- $11\alpha,15S$ -dihydroxy-prosta-Formal Name:

5Z,13E-dien-1-oic acid, monosodium salt

Synonyms: Epoprostenol, PGI₂, Prostacyclin

MF: $C_{20}H_{31}O_5 \cdot Na$

FW: 374.5 ≥98% **Purity:**

≥6 months at -20°C Stability: Supplied as: A crystalline solid

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Laboratory Procedures

For long term storage, we suggest that prostaglandin I2 (sodium salt) (PGI3) be stored as supplied at -20°C. It should be stable for at least six months.

PGI2 is a hygroscopic crystalline solid soluble in water. It is unstable at neutral or acidic pH. On exposure to open air, the compound will absorb moisture and hydrolyze rapidly to 6-keto $PGF_{1\alpha}$. An aqueous stock solution of PGI_2 can be prepared by dissolving the crystalline solid directly in basic buffers (pH >10.2). The solubility of PGI2 in PBS (pH 9.0) is approximately 11 mg/ml. Solutions of PGI₂ at physiologic pH and room temperature will have a half-life from 1 to 12 minutes depending on buffer concentration.^{1,2}

 PGI_2 is an unstable cyclooxygenase metabolite detected first in vascular endothelial cells. 1,3,4 It elevates platelet cAMP and is a potent vasodilator and inhibitor of human platelet aggregation with an IC $_{50}$ value of 5 nM.⁵ PGI $_{2}$ is stable in basic buffers (pH = 8), but it is rapidly hydrolyzed to 6-keto $PGF_{1\alpha}$ in neutral or acidic solutions. The half-life is short both in vivo and in vitro, ranging from 30 seconds to a few minutes. PGI2 is administered by continuous infusion in humans for the treatment of idiopathic pulmonary hypertension.⁶

References

- 1. Stehle, R.G. Physical chemistry, stability, and handling of prostaglandins E2, F20, D2 and I2: A critical summary. Methods Enzymol. 86, 436-459 (1982).
- Moncada, S. Biology and therapeutic potential of prostacyclin. Stroke 14, 157-168 (1983).
- Moncada, S., Gryglewski, R., Bunting, S., et al. An enzyme isolated from arteries transforms prostaglandin endoperoxides to an unstable substance that inhibits platelet aggregation. Nature 263, 663-665 (1976).
- Johnson, R.A., Morton, D.R., Kinner, J.H., et al. The chemical structure of prostaglandin X (prostacyclin). Prostaglandins 12, 915-928 (1976).
- 5. Aristoff, P.A., Johnson, P.D., and Harrison, A.W. Synthesis of 9-substituted carbacyclin analogues. J. Org. Chem. 48, 5341-5348 (1983).
- McLaughlin, V.V., Genthner, D.E., Panella, M.M., et al. Reduction in pulmonary vascular resistance with long-term epoprostenol (prostacyclin) therapy in primary pulmonary hypertension. N. Engl. J. Med. 338, 273-277 (1998).

Related Products

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

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