

# Limaprost

## Catalog No: tcsc2912



### Available Sizes

**Size:** 2mg

**Size:** 5mg

**Size:** 10mg

**Size:** 50mg



### Specifications

**CAS No:**

74397-12-9

**Formula:**

$C_{22}H_{36}O_5$

**Pathway:**

Immunology/Inflammation

**Target:**

PGE synthase

**Purity / Grade:**

>98%

**Solubility:**

DMSO :  $\geq$  40 mg/mL (105.12 mM)

**Alternative Names:**

17 $\alpha$ ,20-dimethyl- $\delta$ 2-PGE1;ONO1206;OP1206

**Observed Molecular Weight:**

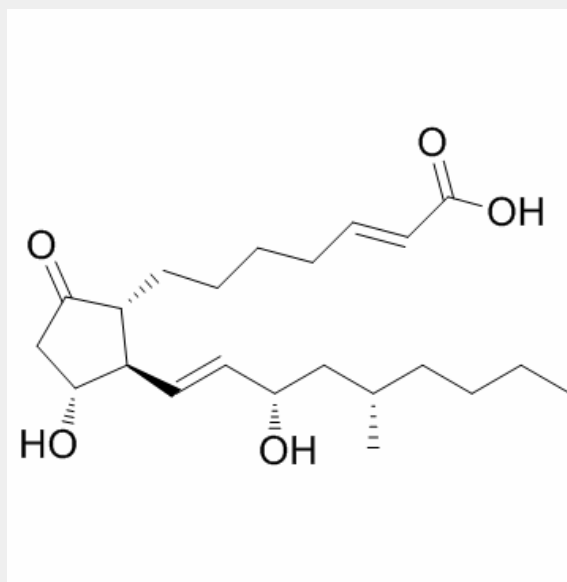
380.52

## Product Description

Limaprost(OP1206) is a PGE1 analog and potent platelet adhesion inhibitor.

Target: Others

Limaprost, an alprostadiol (prostaglandin E1) analogue, is a vasodilator that increases blood flow and inhibits platelet aggregation. Limaprost is a n analog of PGE1 with structural modifications intended to give a prolonged half-life and greater potency. It is orally active in both guinea pigs and rats at doses of 100 mg/kg as an inhibitor of ADP and collagen induced platelet aggregation. It is 10-1,000 times more potent than PGE1 as a platelet adhesive inhibitor, measured in vitro. Intra-coronary injection (100 ng/kg) or intravenous injection (3 mg/kg) in anesthetized dogs causes vasodilation and increased coronary blood flow by 60-80%. Significant hypotensive effects were seen at 100 and 300 mg/kg orally in rats [1].



All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!