PRODUCT INFORMATION



COOL

(+)-15-epi Cloprostenol

Item No. 10006692

CAS Registry No.: 54276-22-1

Formal Name: (+)-9a,11a,15S-trihydroxy-16-(3-

chlorophenoxy)-17,18,19,20-tetranor-

prosta-5Z,13E-dien-1-oic acid

Synonyms: D-Cloprostenol, (+)-15(S)-Cloprostenol

MF: C22H29CIO6 FW: 424.9 **Purity:**

UV/Vis.: λ_{max} : 204, 275 nm A solution in ethanol Supplied as:

-20°C Storage: Stability: ≥2 years

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



(+)-15-epi Cloprostenol is supplied as a solution in ethanol. To change the solvent, simply evaporate the ethanol under a gentle stream of nitrogen and immediately add the solvent of choice. Solvents such as DMSO and dimethyl formamide purged with an inert gas can be used. The solubility of (+)-15-epi cloprostenol in these solvents is approximately 50 mg/ml.

(+)-15-epi Cloprostenol is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, the ethanolic solution of (+)-15-epi cloprostenol should be diluted with the aqueous buffer of choice. The solubility of (+)-15-epi cloprostenol in PBS (pH 7.2) is approximately 1 mg/ml. We do not recommend storing the aqueous solution for more than one day.

Description

Cloprostenol is a synthetic prostaglandin $F_{2\alpha}$ (PGF_{2 α}) analog and a potent FP receptor agonist. (+)-15-epi Cloprostenol is the 15(S), or 15 β -hydroxy enantiomer of (+)-cloprostenol. This epimer is less active by several orders of magnitude as an FP receptor ligand when compared to 15(R)-cloprostenol. However, the specific activity of this isomer has not been well studied.

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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