PRODUCT INFORMATION



Klotho-derived Peptide 1 (56-87) (human) (trifluoroacetate salt)

Item No. 38166

Synonym: KP1 (56-87)

Peptide Sequence: FQGTFPDGFLWAVGSAAYQTEGGWQQHGKG-OH H-Phe-Gln-Gly-Thr-Phe-Pro-Asp-Gly-Phe-Leu-

MF: C₁₄₉H₂₀₃N₃₉O₄₃ • XCF₃COOH

Trp - Ala - Val - Gly - Ser - Ala - Ala - Tyr - Gln - Thr -FW: 3,228.4 Glu-Gly-Gly-Trp-Gln-Gln-His-Gly-Lys-Gly-OH**Purity:** ≥95%

Supplied as: A solid • XCF₃COOH -20°C Storage:

Stability: ≥4 years

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

Laboratory Procedures

Klotho-derived peptide 1 (KP1) (56-87) (human) (trifluoroacetate salt) is supplied as a solid. A stock solution may be made by dissolving the KP1 (56-87) (human) (trifluoroacetate salt) in water. We do not recommend storing the aqueous solution for more than one day.

Description

KP1 (56-87) is a peptide derived from human Klotho protein, which has roles in disrupting TGF-β signaling.¹ It binds to TGF-β receptor type 1 (TGFBR2), and TGF-β receptor type 2 (TGFBR2; K₄s = 1.41 and 14.6 μM, respectively). Preincubation with KP1 (10 μg/ml) inhibits TGF-β-induced increases in fibronectin and α-smooth muscle actin (α-SMA) levels in NRK-49F rat fibroblasts. In vivo, KP1 (1 mg/kg per day) selectively localizes to damaged kidneys and reduces serum creatine and blood urea nitrogen levels, markers of kidney function, as well as reduces kidney fibrosis, in mouse models of unilateral ureteral obstruction (UUO) and unilateral ischemia-reperfusion injury-induced renal fibrosis.

Reference

1. Yuan, Q., Ren, Q., Li, L., et al. A Klotho-derived peptide protects against kidney fibrosis by targeting TGF-β signaling. Nat. Commun. 13(1), 438 (2022).

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.

WARRANTY AND LIMITATION OF REMEDY

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