

Lexacalcitol

Catalog No: tcsc0399



Available Sizes

Size: 1mg

Size: 5mg

Size: 10mg

Size: 25mg



Specifications

CAS No:

131875-08-6

Formula:

$C_{29}H_{48}O_4$

Pathway:

Vitamin D Related

Target:

VD/VDR

Purity / Grade:

>98%

Solubility:

10 mM in DMSO

Alternative Names:

KH 106;KH 1060

Observed Molecular Weight:

460.69

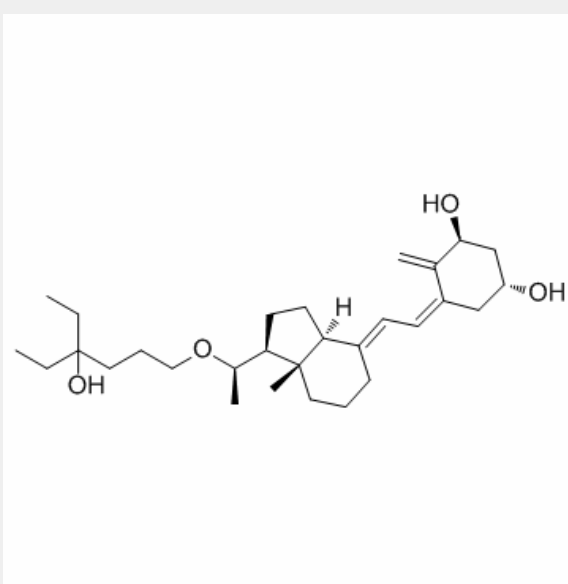
Product Description

Lexacalcitol (KH1060) is over 100 times more active than 1 α ,25-dihydroxyvitamin D₃ [1 α ,25-(OH)₂D₃], as judged by in vitro antiproliferative and cell differentiating assays.

IC₅₀ value:

Target:

KH1060 metabolism could be blocked by the cytochrome P450 inhibitor, ketoconazole. KH1060 was not an effective competitor of C₂₄ oxidation of 1 α ,25-(OH)₂D₃. Certain hydroxylated metabolites of KH1060 retained significant biological activity in vitamin D-dependent reporter gene systems (chloramphenicol acetyltransferase). KH 1060 inhibited PBMC proliferation and decreased TNF- α levels in IBD patients and this effect was synergistic with anti-TNF- α . VDR protein levels were significantly increased by PBMC treatment with KH 1060 or anti-TNF- α or their combination in ulcerative colitis (UC) patients, and decreased in Crohn's disease (CD) patients, treating the cells with KH 1060. A synergistic inhibition was registered combining KH 1060 and anti-TNF, at well-defined concentrations. 0.1 nM KH 1060 produced a significant decrease in TNF- α levels, determined by ELISA, although less remarkable than in the presence of anti-TNF.



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