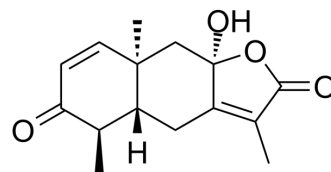


Eudebeiolide B

Cat. No.:	HY-N11775
CAS No.:	1934299-51-0
Molecular Formula:	C ₁₅ H ₁₈ O ₄
Molecular Weight:	262.3
Target:	NF-κB; Akt; Nuclear Factor of activated T Cells (NFAT); Phosphatase
Pathway:	NF-κB; PI3K/Akt/mTOR; Immunology/Inflammation; Metabolic Enzyme/Protease
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Eudebeiolide B is a compound that can be isolated from <i>Salvia plebeia</i> R. Br. Eudebeiolide B inhibits osteoclastogenesis by regulating RANKL-induced NF-κB, c-Fos and calcium signaling. Eudebeiolide B can be used for osteoclast-related diseases research ^[1] .																	
IC₅₀ & Target	Akt	NF-κB																
In Vitro	<p>Eudebeiolide B (1-30 μM, 1 hour) suppresses RANKL-induced osteoclast differentiation and function in mouse bone marrow macrophages (BMMs)^[1].</p> <p>Eudebeiolide B (1-30 μM, 1 hour) inhibits the expression of osteoclastogenesis-related marker genes and RANKL-mediated cellular signaling^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Differentiation Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>Bone marrow macrophages (BMMs)</td> </tr> <tr> <td>Concentration:</td> <td>1-30 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>1 hours</td> </tr> <tr> <td>Result:</td> <td>Inhibited RANKL-induced osteoclast differentiation of BMMs, bone resorption, and promotes osteoblast differentiation.</td> </tr> </table> <p>RT-PCR^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>Bone marrow macrophages (BMMs), MC3T3-E1 cells</td> </tr> <tr> <td>Concentration:</td> <td>10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>1 hours</td> </tr> <tr> <td>Result:</td> <td>Downregulated the expression of NFATc1 and c-fos, transcription factors induced by RANKL. Attenuated the RANKL-induced expression of osteoclastogenesis-related genes, including Ctsk, MMP9 and DC-STAMP. Induced the expression of alkaline phosphatase (ALP) and calcium accumulation during</td> </tr> </table>		Cell Line:	Bone marrow macrophages (BMMs)	Concentration:	1-30 μM	Incubation Time:	1 hours	Result:	Inhibited RANKL-induced osteoclast differentiation of BMMs, bone resorption, and promotes osteoblast differentiation.	Cell Line:	Bone marrow macrophages (BMMs), MC3T3-E1 cells	Concentration:	10 μM	Incubation Time:	1 hours	Result:	Downregulated the expression of NFATc1 and c-fos, transcription factors induced by RANKL. Attenuated the RANKL-induced expression of osteoclastogenesis-related genes, including Ctsk, MMP9 and DC-STAMP. Induced the expression of alkaline phosphatase (ALP) and calcium accumulation during
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	MC3T3-E1 osteoblast differentiation.
Western Blot Analysis ^[1]	
Cell Line:	Bone marrow macrophages (BMMs)
Concentration:	10 μ M
Incubation Time:	1 hours
Result:	Inhibited the phosphorylation of Akt and NF- κ B p65. Downregulated the expression of CREB, Btk and phospholipase PLC γ 2 in RANKL-induced calcium signaling.
In Vivo	Eudebeiolide B (5 or 10 mg/kg, i.g., once daily for 6 weeks) prevents OVX-induced bone loss in an ovariectomized (OVX) mouse model ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
Animal Model:	Ovariectomized (OVX) mouse model ^[1]
Dosage:	5 or 10 mg/kg
Administration:	After 6 weeks of ovarian resection, intragastric injection (i.g.) once daily for 6 weeks.
Result:	Prevented bone mineral density (BMD) loss and bone mineral content (BMC) loss compared to the OVX mice.

REFERENCES

[1]. Kim MH, et al. Eudebeiolide B Inhibits Osteoclastogenesis and Prevents Ovariectomy-Induced Bone Loss by Regulating RANKL-Induced NF- κ B, c-Fos and Calcium Signaling. *Pharmaceuticals (Basel)*. 2020 Dec 16;13(12):468.

Caution: Product has not been fully validated for medical applications. For research use only.

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