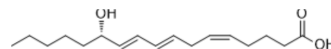


## 12S-HHT

Cat. No.:	HY-113330
CAS No.:	54397-84-1
Molecular Formula:	C <sub>17</sub> H <sub>28</sub> O <sub>3</sub>
Molecular Weight:	280.4
Target:	Leukotriene Receptor; Endogenous Metabolite
Pathway:	GPCR/G Protein; Metabolic Enzyme/Protease
Storage:	Solution, -20°C, 2 years



### BIOLOGICAL ACTIVITY

<b>Description</b>	12S-HHT (12(S)-HHTre) is an enzymatic product of prostaglandin H <sub>2</sub> (PGH <sub>2</sub> ) derived from cyclooxygenase (COX)-mediated arachidonic acid metabolism. 12S-HHT is an endogenous ligand for BLT2 that fully activates BLT2 in vivo. 12S-HHT suppresses UV-induced IL-6 synthesis in keratinocytes, exerting an anti-inflammatory activity <sup>[1][2]</sup> .									
<b>IC<sub>50</sub> &amp; Target</b>	Human Endogenous Metabolite	BLT2								
<b>In Vitro</b>	<p>12S-HHT (0-150 nM; 3 hours) has anti-inflammatory activity by attenuating the UVB-induced IL-6 synthesis in HaCaT cells<sup>[2]</sup>. 12S-HHT inhibits the UVB-stimulated p38 MAPK/NF-κB pathway by up-regulating MKP-1, which leads to the suppression of IL-6 synthesis<sup>[2]</sup>.</p> <p>12S-HHT is an endogenous agonist for BLT2<sup>[3]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Western Blot Analysis<sup>[2]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>HaCaT cells</td> </tr> <tr> <td>Concentration:</td> <td>0, 12.5, 25, 75 or 150 nM</td> </tr> <tr> <td>Incubation Time:</td> <td>3 hours</td> </tr> <tr> <td>Result:</td> <td>UVB (5 mJ/cm<sup>2</sup>) irradiation markedly up-regulated IL-6 synthesis and release, which was suppressed by the treatment with 12-HHT in a concentration-dependent manner.</td> </tr> </table>		Cell Line:	HaCaT cells	Concentration:	0, 12.5, 25, 75 or 150 nM	Incubation Time:	3 hours	Result:	UVB (5 mJ/cm <sup>2</sup> ) irradiation markedly up-regulated IL-6 synthesis and release, which was suppressed by the treatment with 12-HHT in a concentration-dependent manner.
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### CUSTOMER VALIDATION

- Cell Rep Med. 2023 May 24;101061.

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### REFERENCES

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[1]. Saeki K, et al. Identification, signaling, and functions of LTB4 receptors. *Semin Immunol.* 2017;33:30-36.

[2]. Lee JW, et al. 12(S)-Hydroxyheptadeca-5Z,8E,10E-trienoic acid suppresses UV-induced IL-6 synthesis in keratinocytes, exerting an anti-inflammatory activity. *Exp Mol Med.* 2012;44(6):378-386.

[3]. Okuno T, et al. Metabolism and biological functions of 12(S)-hydroxyheptadeca-5Z,8E,10E-trienoic acid. *Prostaglandins Other Lipid Mediat.* 2021;152:106502.

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

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