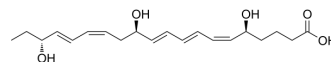


## Resolvin E1

Cat. No.:	HY-114041
CAS No.:	552830-51-0
Molecular Formula:	C <sub>20</sub> H <sub>30</sub> O <sub>5</sub>
Molecular Weight:	350.45
Target:	Endogenous Metabolite
Pathway:	Metabolic Enzyme/Protease
Storage:	Solution, -20°C, 2 years



### BIOLOGICAL ACTIVITY

<b>Description</b>	Resolvin E1 (RvE1), a potent endogenous pro-resolving mediator of inflammation, is derived from omega-3 fatty acid eicosapentaenoic acid (EPA). Resolvin E1 is endogenously biosynthesized from EPA in the presence of Aspirin during the spontaneous resolution phase of acute inflammation, where specific cell-cell interactions occur. Resolvin E1 possesses unique counterregulatory actions that inhibit polymorphonuclear leukocyte (PMN) transendothelial migration. Resolvin E1 also acts as a potent inhibitor of leukocyte infiltration, dendritic cell migration, and IL-12 production <sup>[1][2]</sup> .								
<b>IC<sub>50</sub> &amp; Target</b>	Human Endogenous Metabolite								
<b>In Vitro</b>	Resolvin E1 (0.1-100 nM) gives concentration-dependent inhibition of TNF-α-induced NF-κB activation with an EC <sub>50</sub> of 1.0 nM in ChemR23-transfected cells. Resolvin E1 (100 nM) increases phosphorylation of extracellular signal-regulated kinase (ERK) mitogen-activated protein (MAP) kinase both in peripheral blood monocytes and HEK-ChemR23 cells <sup>[2]</sup> . Resolvin E1 specifically interacts with the LTB <sub>4</sub> receptor BLT1 on neutrophils and ChemR23 on monocytes to regulate leukocytes during inflammation. Resolvin E1 also stimulates the uptake and clearance of local cytokine <sup>[3]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.								
<b>In Vivo</b>	Resolvin E1 (1.0 µg per mouse; 50 µg/kg; i.p.; on days -8, -1, and 0) protects mice from TNBS (2,4,6-trinitrobenzene sulfonic acid) colitis <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.								
	<table border="1"> <tr> <td>Animal Model:</td> <td>6- to 8-week-old female BALB/c mice (TNBS colitis model)<sup>[1]</sup></td> </tr> <tr> <td>Dosage:</td> <td>1.0 µg per mouse; 50 µg/kg</td> </tr> <tr> <td>Administration:</td> <td>i.p.; on days -8, -1, and 0 before the induction of colitis</td> </tr> <tr> <td>Result:</td> <td>Reduced overall mortality, which was 25% and 62.5% at 1.0 µg per mouse and 50 µg/kg, respectively.</td> </tr> </table>	Animal Model:	6- to 8-week-old female BALB/c mice (TNBS colitis model) <sup>[1]</sup>	Dosage:	1.0 µg per mouse; 50 µg/kg	Administration:	i.p.; on days -8, -1, and 0 before the induction of colitis	Result:	Reduced overall mortality, which was 25% and 62.5% at 1.0 µg per mouse and 50 µg/kg, respectively.
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### REFERENCES

[1]. Arita M, et al. Resolvin E1, an endogenous lipid mediator derived from omega-3 eicosapentaenoic acid, protects against 2,4,6-trinitrobenzene sulfonic acid-induced colitis. Proc Natl Acad Sci U S A. 2005 May 24;102(21):7671-6.

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[2]. Schwab JM, et al. Resolvin E1 and protectin D1 activate inflammation-resolution programmes. Nature. 2007 Jun 14;447(7146):869-74.

[3]. Hasturk H, et al. Resolvin E1 regulates inflammation at the cellular and tissue level and restores tissuehomeostasis in vivo. J Immunol. 2007 Nov 15;179(10):7021-9.

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

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