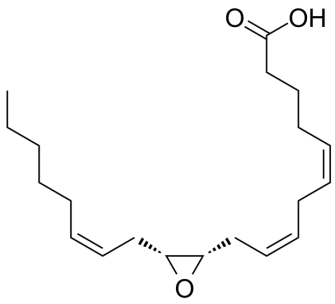
(±)11(12)-EET

Cat. No.:	HY-130494
CAS No.:	123931-40-8
Molecular Formula:	$C_{20}H_{32}O_3$
Molecular Weight:	320.47
Target:	NOD-like Receptor (NLR)
Pathway:	Immunology/Inflammation
Storage:	Solution, -20°C, 2 years



BIOLOGICAL ACTIVITY

Description

(±)11(12)-EET is a NLRP3 inflammasome inhibitor. (±)11(12)-EET can be used for the research of anti-inflammatory, angiogenic and cardioprotective[1][2][3][4][6].

IC_{50} & Target

NLRP3 inflammasome

In Vitro

(±)11(12)-EET (5 μM; 10 mimutes; macrophages) depresses NLRP3 protein expression, dramatically decreases the expression of pro-IL-1β in cells and the supernatant and reduces the intracellular ROS[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Western Blot Analysis[1]

Cell Line:	Macrophages
Concentration:	5 μM
Incubation Time:	10 mimutes
Result:	Depresses NLRP3 protein expression.

Immunofluorescence[1]

Cell Line:	Macrophages
Concentration:	5 μM
Incubation Time:	10 mimutes
Result:	Reduced the intracellular ROS.

In Vivo

(±)11(12)-EET increases adhesion of isolated peripheral blood leukocytes in a chamber coated with P-selectin and ICAM-1 in 50 μg/kg[5].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Luo XQ, et al. Epoxyeicosatrienoic acids inhibit the activation of NLRP3 inflammasome in murine macrophages. J Cell Physiol. 2020;235(12):9910-9921.

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[4]. Capdevila JH, et al. Cytochrome P450 and arachidonic acid bioactivation. Molecular and functional properties of the arachidonate monooxygenase. J Lipid Res. 2000;41(2):163-181.

[5]. Wang Z, et al. Arachidonic acid inhibits basolateral K channels in the cortical collecting duct via cytochrome P-450 epoxygenase-dependent metabolic pathways. Am J Physiol Renal Physiol. 2008;294(6):F1441-F1447.

[6]. Spector AA. Arachidonic acid cytochrome P450 epoxygenase pathway. J Lipid Res. 2009;50 Suppl(Suppl):S52-S56.

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