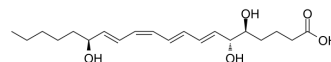


Lipoxin A4

Cat. No.:	HY-113509
CAS No.:	89663-86-5
Molecular Formula:	C ₂₀ H ₃₂ O ₅
Molecular Weight:	352.47
Target:	Interleukin Related; Endogenous Metabolite
Pathway:	Immunology/Inflammation; Metabolic Enzyme/Protease
Storage:	Solution, -20°C, 2 years



BIOLOGICAL ACTIVITY

Description	Lipoxin A4 (LXA4), an endogenous lipoxygenase-derived eicosanoid mediator, has potent dual pro-resolving and anti-inflammatory properties ^[1] . Lipoxin A4 inhibits proliferation and inflammatory cytokine/chemokine production of human epidermal keratinocytes (NHEKs) associated with the ERK1/2 and NF-κB pathways ^[2] . Lipoxin A4 inhibits serum amyloid A (SAA)-mediated IL-8 release with an IC ₅₀ value of 25.74 nM ^[3] .															
IC₅₀ & Target	Human Endogenous Metabolite															
In Vitro	<p>Lipoxin A4 (LXA4) inhibits the expression of IL-6 and IL-8 in NHEKs^[2]. Lipoxin A4 downregulates the expression of cyclin D1^[2]. Lipoxin A4 also suppresses the ERK1/2 phosphorylation and NF-κB-p65 nuclear translocation of NHEKs^[2]. LXA4 (100 nM; preincubation for 30 minutes) inhibits the proliferation of NHEKs with or without stimulating by LPS (10 μg/mL)^[2]. LXA4 pretreatment (100 nM for 30 minutes) downregulates the LPS-induced secretion and expression of HMGB1 in keratinocytes^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Proliferation Assay^[2]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>Normal human epidermal keratinocytes (NHEKs)</td> </tr> <tr> <td>Concentration:</td> <td>100 nM</td> </tr> <tr> <td>Incubation Time:</td> <td>30 minutes</td> </tr> <tr> <td>Result:</td> <td>A significant increase in proliferation of NHEKs after 7 days of stimulation with LPS (10 μg/mL) was seen. However, there was a significant decrease in the proliferation of NHEKs when pretreated with LXA4 for 30 min.</td> </tr> </table> <p>Western Blot Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>Normal human epidermal keratinocytes (NHEKs)</td> </tr> <tr> <td>Concentration:</td> <td>100 nM</td> </tr> <tr> <td>Incubation Time:</td> <td>30 minutes.</td> </tr> </table>		Cell Line:	Normal human epidermal keratinocytes (NHEKs)	Concentration:	100 nM	Incubation Time:	30 minutes	Result:	A significant increase in proliferation of NHEKs after 7 days of stimulation with LPS (10 μg/mL) was seen. However, there was a significant decrease in the proliferation of NHEKs when pretreated with LXA4 for 30 min.	Cell Line:	Normal human epidermal keratinocytes (NHEKs)	Concentration:	100 nM	Incubation Time:	30 minutes.
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Result:

HMGB1 protein levels in the cytoplasm of NHEKs were induced by LPS, which were decreased after preincubation with LXA4 but decreased in the nucleus after stimulation with LPS.

CUSTOMER VALIDATION

- Drug Alcohol Depend. 1 August 2022, 109537.

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REFERENCES

- [1]. Feng Hu, et al. Lipoxin A4 inhibits proliferation and inflammatory cytokine/chemokine production of human epidermal keratinocytes associated with the ERK1/2 and NF- κ B pathways. *J Dermatol Sci*. 2015 Jun;78(3):181-8.
- [2]. Steven Bozinovski, et al. Serum amyloid A opposes lipoxin A₄ to mediate glucocorticoid refractory lung inflammation in chronic obstructive pulmonary disease. *Proc Natl Acad Sci U S A*. 2012 Jan 17;109(3):935-40.
- [3]. Xinxin Liu, et al. Lipoxin A4 and its analog suppress inflammation by modulating HMGB1 translocation and expression in psoriasis. *Sci Rep*. 2017 Aug 2;7(1):7100.
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Caution: Product has not been fully validated for medical applications. For research use only.

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