

## Produktinformation



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### SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien T. +43(0)1 489 3961-0 F. +43(0)1 489 3961-7 <u>mail@szabo-scandic.com</u> www.szabo-scandic.com

## **PRODUCT** INFORMATION



### Autotaxin (human, recombinant)

Item No. 10803

#### **Overview and Properties**

Synonyms:	ATX, Ectonucleotide Pyrophosphatase/Phosphodiesterase-2, ENPP-2, Lyso-PLD, Lysophospholipase D
Source:	Active recombinant human autotaxin expressed in insect cells
Amino Acids:	36-863
Uniprot No.:	Q13822-1
Molecular Weight:	98.8 kDa
Storage:	-80°C (as supplied)
Stability:	≥2 years
Purity:	≥90% estimated by SDS-PAGE
Supplied in:	50 mM Tris-HCl, pH 8.0, with 150 mM sodium chloride and 20% glycerol
Protein	
Concentration:	<i>batch specific</i> mg/ml
Activity:	<i>batch specific</i> U/ml
Specific Activity:	batch specific U/mg
Unit Definition:	One unit is defined as the amount of enzyme required to produce 1 µmol of <i>p</i> -nitrophenol
	per minute at 37°C in 50 mM Tris-HCl, pH 9.0, containing <i>bis</i> -nitrophenol phosphate.

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

#### Images





Autotaxin activity is determined by measuring hydrolysis of bis-nitrophenol phosphate. Production of *p*-nitrophenol is measured by monitoring the increase in absorbance at 410 nm.

#### WARNING THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

#### SAFETY DATA

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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#### CAYMAN CHEMICAL

1180 EAST ELLSWORTH RD ANN ARBOR, MI 48108 · USA PHONE: [800] 364-9897 [734] 971-3335 FAX: [734] 971-3640 CUSTSERV@CAYMANCHEM.COM WWW.CAYMANCHEM.COM

# **PRODUCT** INFORMATION



#### Description

Autotaxin, also known as ectonucleotide pyrophosphatase/phosphodiesterase 2 (ENPP2), is a secreted lysophospholipase D (lysoPLD) that hydrolyzes lysophosphatidylcholine (LPC) to lysophosphatidic acid (LPA), which has roles in cell proliferation, survival, and migration.<sup>1</sup> It is synthesized as a zymogen and is composed of a N-terminal signal peptide, which is cleaved during maturation, two somatomedin B-like domains, a phosphodiesterase (PDE) catalytic domain, and a C-terminal nuclease-like domain.<sup>2,3</sup> It is constitutively active and inhibited by LPA in a negative feedback loop.<sup>4,5</sup> Autotaxin is secreted primarily by adipose tissue and endothelial cells and has been found in numerous biological fluids, including the blood, cerebrospinal fluid, and saliva. It is overexpressed in a variety of cancers, including glioblastoma multiforme, melanoma, and hepatocarcinoma.<sup>4-6</sup> Autotaxin has roles in cell motility, immune regulation, and embryogenesis.<sup>4,5</sup> Knockout of *Enpp2* is embryonic lethal in mice.<sup>4</sup> Autotaxin activity is increased in the serum of patients with a variety of conditions, including rheumatoid arthritis, chronic hepatitis C, or cholestasis, and serum autotaxin levels are increased in patients with asthma, acute respiratory distress syndrome (ARDS), or coronavirus disease 2019 (COVID-19).<sup>3,7</sup> Formulations containing autotaxin inhibitors have been used in clinical trials for the treatment of various diseases, including idiopathic pulmonary fibrosis (IPF), metabolic disorders, and cancer.<sup>7</sup> Cayman's Autotaxin (human, recombinant) protein can be used for enzyme activity assays.

#### References

- 1. Nishimasu, H., Okudaira, S., Hama, K., *et al.* Crystal structure of autotaxin and insight into GPCR activation by lipid mediators. *Nat. Struct. Mol. Biol.* **18(2)**, 205-12 (2011).
- Jansen, S., Stefan, C., Creemers, J.W.M., et al. Proteolytic maturation and activation of autotaxin (NPP2), a secreted metastasis-enhancing lysophospholipase D. J. Cell Sci. 118(14), 3081-9 (2011).
- 3. Magkrioti, C., Galaris, A., Kanellopoulou, P., *et al.* Autotaxin and chronic inflammatory diseases. *J. Autoimmun.* **104**, 102327 (2019).
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- Ramesh, S., Govindarajulu, M., Suppiramaniam, V., et al. Autotaxin-lysophosphatidic acid signaling in Alzheimer's disease. Int. J. Mol. Sci. 19(7), 1827 (2018).
- 7. Ntatsoulis, K., Karampitsakos, T., Tsitoura, E., *et al.* Commonalities between ARDS, pulmonary fibrosis and COVID-19: The potential of autotaxin as a therapeutic target. *Front. Immunol.* **12**, 687397 (2021).

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