

AG-20B-0064

13-Feb-2014

anti-IL-1 α (p18) (mouse), mAb (Teo-1)
[Interleukin-1 α]

AG-20B-0064-C100	100 μ g
Clone	Teo-1
Source/Host	Purified from concentrated hybridoma tissue culture supernatant.
Isotype	Mouse IgG
Immunogen	Recombinant mouse mature IL-1 α .

Handling / Storage

Shipping	BLUE ICE
Short Term Storage	+4°C
Long Term Storage	-20°C

After opening, prepare aliquots and store at -20°C. Avoid freeze/thaw cycles.

Use / Stability

Stable for at least 1 year after receipt when stored at -20°C.

MSDS available at www.adipogen.com or upon request.

Product Specifications

Specificity	Recognizes mouse IL-1 α p18 cleaved and full-length fragments.
Species Crossreactivity	Mouse
Application	Western Blot: (1 μ g/ml) ELISA
Purity	\geq 95% (SDS-PAGE)
Formulation	Liquid. In PBS containing 10% glycerol and 0.02% sodium azide.
Concentration	1mg/ml

Product Description

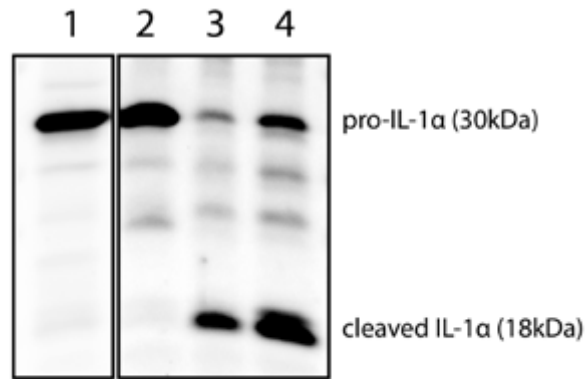
The most prominent members of the interleukin-1 (IL-1) superfamily are IL-1 α and IL-1 β . They lack a signal peptide and are secreted by an unconventional, endoplasmic reticulum-Golgi-independent mechanism. IL-1 α was reported to be more widely and constitutively expressed and has intracellular functions, but also acts locally in a membrane-bound form by activating IL-1R1. Additionally, passive release of IL-1 α upon cell death can trigger a sterile inflammatory response to dying cells. The cleavage of IL-1 α is not mediated by caspase-1 and is not required for binding to IL-1R1. Recently it has been observed that all activators of the inflammasome NLRP3/NALP3 induce the simultaneous secretion of IL-1 α and IL-1 β . Although most activators fully rely on the inflammasome for IL-1 α secretion, some induce the processing and secretion of IL-1 α in an inflammasome-independent manner.

WARNING: Intended for research use only. This product is not intended or approved for human, diagnostics, therapeutic or veterinary use. Use of this product for human or animal testing is extremely hazardous and may result in disease, severe injury, or death. **MATERIAL SAFETY DATA:** Review the complete Material Safety Data Sheet before use.

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- 1: Lysate of LPS-primed BMDCs
- 2: Supernatant of LPS +ATP treated BMDCs
- 3: Supernatant of LPS + Nigericin treated BMDCs
- 4: Supernatant of LPS + MSU treated BMDCs

Figure 1: Mouse IL-1α (full-length p30 and cleaved p18 fragments) are detected by immunoblotting using anti-IL-1α (p18) (mouse), mAb (Teo-1) (Prod. No AG-20B-0064).

Method: IL-1α was analyzed by Western blot in cell extracts of bone marrow-derived dendritic cells (BMDCs) treated by LPS and the several inflammasome activators as indicated in the figure. Cell extracts were separated by SDS-PAGE under reducing conditions, transferred to nitrocellulose and incubated with anti-IL-1α (p18) (mouse), mAb (Teo-1) (1μg /ml). After addition of an anti-mouse secondary antibody coupled to HRP, proteins were visualized by a chemiluminescence detection system.

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