

Product datasheet for AM26314PU-N

Aoc3 Rat Monoclonal Antibody [Clone ID: 7-88]

Product data:

OriGene Technologies, Inc.

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Product Type:	Primary Antibodies
Clone Name:	7-88
Applications:	FN, IF, IHC, IP
Recommended Dilution:	 Immunohistochemistry on Frozen Sections (Ref.2,3): Tissue sections were fixed in acetone and incubated with antibody 7-88 for 20 minutes at room temperature. As negative control an irrelevant isotype-matched antibody was used (Ref.2). Flow Cytometry: Stains the extracellular domain of mouse VAP-1 in CHO cells transfected with mouse VAP-1 cDNA. As positive control anti-VAP-1 clone 7-106 was used and as negative control an isotype-matched control antibody (Ref. 2). Functional Assays (Ref.2,4): Antibody 7-88 (200µg) was intravenously injected which resulted in the inhibition of leukocyte trafficking in inflamed peritoneum (Ref.2). Immunofluorescence (Ref.5). Immunoprecipitation (Ref.1). Positive Control: Mouse VAP-1-transfected CHO cells (Ref.2). Negative Control: Mock-transfected CHO cells (Ref.2).
Reactivity:	Mouse
Host:	Rat
lsotype:	lgG2b
Clonality:	Monoclonal
Immunogen:	Vessels from mouse lymph nodes
Specificity:	The monoclonal antibody 7-88 recognizes mouse Vascular Adhesion Protein-1 (VAP-1) which is a glycosylated homodimeric membrane protein consisting of two 90 kDa subunits connected by disulfide bonds. It inhibits migration of granulocytes and monocytes in acute models of inflammation.
Formulation:	PBS State: Purified State: Liquid 0.2 μm filtered lg fraction Stabilizer: 0.1% BSA
Concentration:	lot specific
Purification:	Protein G Chromatography



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	Aoc3 Rat Monoclonal Antibody [Clone ID: 7-88] – AM26314PU-N
Conjugation:	Unconjugated
Storage:	Store undiluted at 2-8°C.
Stability:	Shelf life: one year from despatch.
Gene Name:	amine oxidase, copper containing 3
Database Link:	Entrez Gene 11754 Mouse O70423
Background:	VAP-1 is a glycosylated homodimeric membrane protein consisting of two 90 kDa subunits connected by disulfide bonds. It contains a short N-terminal cytoplasmic tail, a single membrane-spanning domain and a large extracellular part. A soluble form of VAP-1 (sVAP-1) has been described, which presumably results from the proteolytic cleavage of membrane- bound VAP-1. Structurally VAP-1 belongs to enzymes called semicarbamizide-sensitive amine oxidases, which contain copper as a cofactor. These enzymes deaminate primary amines in a reaction producing hydrogen peroxide, aldehyde, and ammonia. VAP-1 is expressed in endothelial cells, smooth muscle cells, adipocytes, and in follicular dendritic cells. In endothelial cells the majority of VAP-1 is stored within intracellular granules and translocated to the surface upon inflammation where it regulates leukocyte tissue infiltration. Furthermore, the end-products formed by VAP-1 can also regulate leukocyte migration by signaling effects, have insulin-like effects in energy metabolism, and can cause vascular damage by direct cytotoxicity. In white adipose tissue of obese and diabetic db-/- mice increased expression of VAP-1 has been observed suggesting that it contributes to the arthrosclerosis and vascular dysfunction observed in these diseases. Moreover, inhibition of VAP-1reduced the accumulation of myeloid cells into tumors and attenuates tumor growth.
Synonyms:	HPAO

Product images:



Frozen Sections: VAP-1 expression in mouse muscle tissue.

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