

OriGene Technologies, Inc.

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Product datasheet for TA326650

ARTS1 (ERAP1) Rabbit Polyclonal Antibody

Product data:

Product Type:	Primary Antibodies
Applications:	IF, IHC, WB
Recommended Dilution:	WB: 1 - 2 ug/mL
Reactivity:	Human, Mouse
Host:	Rabbit
lsotype:	IgG
Clonality:	Polyclonal
Immunogen:	ERAP1 antibody was raised against a 19 amino acid peptide near the carboxy terminus of human ERAP.
Formulation:	ERAP1 Antibody is supplied in PBS containing 0.02% sodium azide.
Concentration:	1 mg/ml
Purification:	ERAP1 Antibody is affinity chromatography purified via peptide column.
Conjugation:	Unconjugated
Storage:	Store at -20°C as received.
Stability:	Stable for 12 months from date of receipt.
Predicted Protein Size:	Predicted: 104 kDa; Observed: 105 kDa
Gene Name:	endoplasmic reticulum aminopeptidase 1
Database Link:	<u>NP_057526</u> <u>Entrez Gene 80898 MouseEntrez Gene 51752 Human</u> <u>Q9NZ08</u>
Background:	The endoplasmic reticulum (ER) aminopeptidase 1 (ERAP1), a member of the peptidase M1 family, plays a central role in peptide trimming, a step required for the generation of most HLA class I-binding peptides (1,2). It is also designated as adipocyte-derived leucine aminopeptidase (A-LAP), puromycin-insensitive leucine-specific aminopeptidase (PILS-AP) and aminopeptidase regulator of TNFR1 shedding (ARTS-1) (3). ERAP1 is localized to the lumen of the ER and induced by interferon. It may be involved in the regulation of blood pressure through the inactivation of angiotensin II and/or the generation of bradykinin in the kidney (3,4).



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ARTS1 (ERAP1) Rabbit Polyclonal Antibody – TA326650 Synonyms: A-LAP; ALAP; APPILS; ARTS-1; ARTS1; ERAAP; ERAAP1; PILS-AP; PILSAP

Protein Families: Druggable Genome, Protease, Secreted Protein

Product images:



Western blot analysis of ERAP1 in SK-N-SH cell lysate with ERAP1 antibody at (A) 1 and (B) 2 ug/ml.

Immunohistochemistry of ERAP1 in mouse brain tissue with ERAP1 antibody at 5 ug/mL.

Immunofluorescence of ERAP1 in mouse brain tissue with ERAP1 antibody at 20 ug/mL.

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