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anti-ATAD5 antibody (Internal Region)

Images



Overview

Quantity:	100 μg
Target:	ATAD5
Binding Specificity:	Internal Region
Reactivity:	Human
Host:	Goat
Clonality:	Polyclonal
Conjugate:	This ATAD5 antibody is un-conjugated
Application:	ELISA, Flow Cytometry (FACS), Immunofluorescence (IF)

Product Details

Purpose:	ATAD5 / FRAG1
Immunogen:	Peptide with sequence C-DNKIYPKNTKKKRVD, from the internal region of the protein sequence according to NP_079133.3.
Sequence:	DNKIYPKNTK KKRVD
Isotype:	IgG
Cross-Reactivity:	Human
Purification:	Purified from goat serum by ammonium sulphate precipitation followed by antigen affinity chromatography using the immunizing peptide.
Grade:	Verified

Target Details

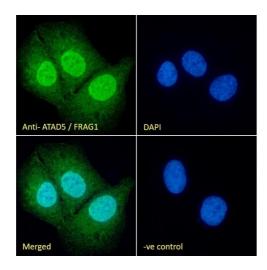
Target:	ATAD5
Alternative Name:	ATAD5 (ATAD5 Products)
Background:	ATAD5, C17orf41, chromosome 17 open reading frame 41, FLJ12735, FRAG1, chromosome fragility associated gene 1, ATPase family, AAA domain containing 5
Gene ID:	79915
NCBI Accession:	NP_079133

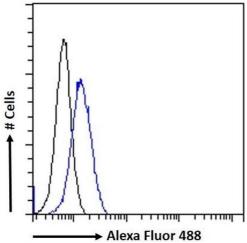
Application Details

Application Notes:	Peptide ELISA: antibody detection limit dilution 1:32000.
Comment:	Immunofluorescence: Strong expression of the protein seen in the nuclei of U2OS cells.
	Recommended concentration: 10µg/ml.
	Flow Cytometry: Flow cytometric analysis of U2OS cells. Recommended concentration:
	10ug/ml.
Restrictions:	For Research Use only

Handling

Format:	Liquid
Concentration:	0.5 mg/mL
Buffer:	Supplied at 0.5 mg/mL in Tris saline, 0.02 % sodium azide, pH 7.3 with 0.5 % bovine serum albumin.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Handling Advice:	Minimize freezing and thawing.
Storage:	-20 °C
Storage Comment:	Aliquot and store at -20°C, with minimal freeze/thawing. A working aliquot may be refrigerated at 4°C for a few weeks and still remain viable.





Immunofluorescence

Image 1. (ABIN190860) Immunofluorescence analysis of paraformaldehyde fixed U2OS cells, permeabilized with 0.15 % Triton. Primary incubation 1hr (10 μ g/mL) followed by Alexa Fluor 488 secondary antibody (2 μ g/mL), showing nuclear staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10 μ g/mL) followed by Alexa Fluor 488 secondary antibody (2 μ g/mL).

Flow Cytometry

Image 2. (ABIN190860) Flow cytometric analysis of paraformaldehyde fixed HeLa cells (blue line), permeabilized with 0.5 % Triton. Primary incubation 1hr (10 μg/mL) followed by Alexa Fluor 488 secondary antibody (1 μg/mL). IgG control: Unimmunized goat IgG (black line) followed by Alexa Fluor 488 secondary antibody.