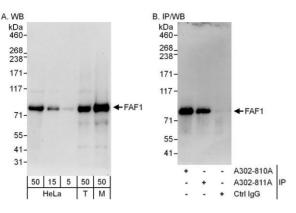
FAF1 Antibody

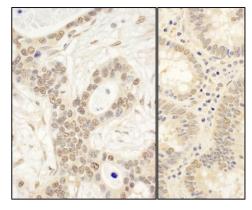
5	,	Protein ID GenelD	NP_008982.1 11124	BETHYL LABORATORIES, INC	
APPLICATIONS	WB, IP, IHC				
SPECIES REACTIVIT	Y Human, Mou	Human, Mouse			
PRESUMED REACTI	VITY Based on 100	Based on 100% sequence identity, this antibody is predicted to react with Rat			
AMOUNT	100 µl	100 µl			
CONCENTRATION	200 µg/ml	200 µg/ml			
STORAGE/SHELF L	FE 2 – 8° C / 1 y	2 - 8° C / 1 year from date of receipt			
PHYSICAL STATE	Liquid	Liquid			
BUFFER	Tris-buffered	Tris-buffered Saline containing 0.1% BSA and 0.09% Sodium Azide			
ISOTYPE	IgG	lgG			
ORIGIN	USA	USA			
PRODUCTION PROCEDURES	Antibody was	Antibody was affinity purified using an epitope specific to FAF1 immobilized on solid support.			
The epitope re Fas (TNFRSF6) 11124).		ecognized by A302-810A maps to a region between residue 25 and 75 of human Associated Factor 1 using the numbering given in entry NP_008982.1 (GeneID centration was determined by extinction coefficient: absorbance at 280 nm of 1.4 of IgG.			
APPLICATIONS	Centrifuge tu	Centrifuge tube to remove product from lid. Optimal working dilutions should be determined experimentally by the investigator. Prepare working dilution immediately before use.			
	Western Blot	1:2	2,000 - 1:10,000		
	Immunoprec	ipitation 2 -	5 µg/mg lysate		
	Immunohisto	,	00 – 1:500. Epitope retriev commended for FFPE tissue	al with citrate buffer pH 6.0 is sections.	
APPLICATION NOT	Goat anti-Ral	Western blot of immunoprecipitates performed using Normal Pig Serum (Cat. No. S100–020), Goat anti–Rabbit Light Chain HRP Conjugate (Cat. No. A120–113P) and 4–8% SDS–PAGE (link to IP–western blot protocol in Additional Info section below).			
		ot of lysates performed using standard western blot reagents and 4–8% SDS-PAGE. noma, Ovarian Carcinoma			
ADDITIONAL INFO	Use the link a	https://www.bethyl.com/product/A302-810A Use the link above to view SDS, a current list of citations, and other product specific information. IP-western blot protocol: https://www.bethyl.com/content/protocol_IP_WB			

This document certifies that this product has met all of the quality control standards defined by Bethyl Laboratories, Inc. Eric McIntush, PhD | Chief Scientific Officer Date: June 21, 2019

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Detection of human and mouse FAF1 by western blot (h&m) and immunoprecipitation (h). *Samples:* Whole cell lysate from HeLa (5, 15 and 50 µg for WB; 1 mg for IP, 20% of IP loaded), HEK293T (T; 50 µg), and mouse NIH 3T3 (M; 50 µg) cells. *Antibodies:* Affinity purified rabbit anti-FAF1 antibody A302-810A used for WB at 0.04 µg/ml (A) and 1 µg/ml (B) and used for IP at 3 µg/mg lysate. FAF1 was also immunoprecipitated by rabbit anti-FAF1 antibody A302-811A, which recognizes a downstream epitope. *Detection:* Chemiluminescence with exposure times of 30 seconds (A) and 10 seconds (B).



Detection of human FAF1 by immunohistochemistry. *Sample:* FFPE section of human ovarian carcinoma (left) and human colon carcinoma (right). *Antibody:* Affinity purified rabbit anti– FAF1 (Cat. No. A302–810A Lot1) used at a dilution of 1:200 (1µg/ml). *Detection:* DAB