

Basic Information

Product Name	Anti-IFT88 Antibody	
Gene Name	IFT88	
Source	Rabbit	
Clonality	Polyclonal	
Isotype	IgG	
Species Reactivity	human, mouse, rat	
Tested Application	WB, IHC, ICC/IF, FCM, ELISA	
Contents	500 ug/ml antibody with PBS, 0.02% NaN3, 1 mg/ml BSA and 50% glycerol.	
Immunogen	E.coli-derived human IFT88 recombinant protein (Position: E198-K814).	
Concentration	500 ug/ml	
Purification	Immunogen affinity purified.	
Observed MW	94 kDa	
Dilution Ratios	Western blot (WB):	1:500-2000
	Immunohistochemistry (IHC):	1:50-400
	Immunocytochemistry/Immunofluorescence (ICC/IF):	1:50-400
	Flow Cytometry (Fixed):	1:50-200
	Enzyme linked immunosorbent assay (ELISA):	1:100-1000
	(Boiling the paraffin sections in 10mM citrate buffer, pH6.0, or PH8.0 EDTA repair liquid for 20 mins is required for the staining of formalin/paraffin sections.) Optimal working dilutions must be determined by end user.	

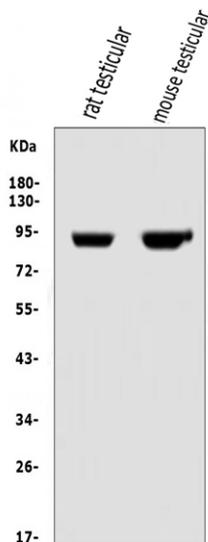
Storage

12 months from date of receipt, -20°C as supplied.

Background Information

Intraflagellar transport protein 88 homolog is a protein that in humans is encoded by the IFT88 gene. It is mapped to 13q12.11. This gene encodes a member of the tetratricopeptide repeat (TPR) family. The encoded protein is involved in cilium biogenesis. Mutations of a similar gene in mouse can cause polycystic kidney disease. Several transcript variants encoding distinct isoforms have been identified for this gene.

Selected Validation Data



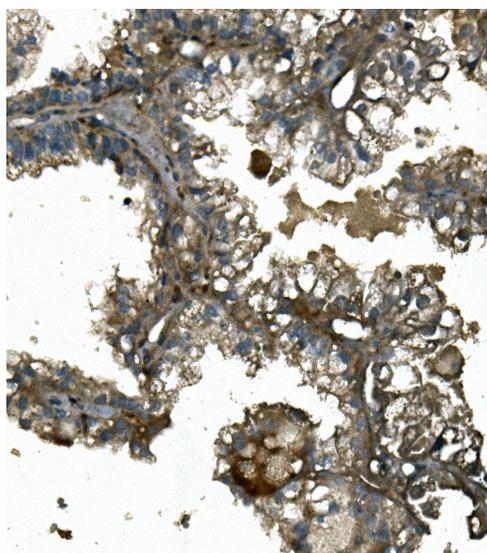
Western blot analysis of IFT88 using anti-IFT88 antibody (A06814-1). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: Rat testicular tissue lysates,

Lane 2: Mouse testicular tissue lysates.

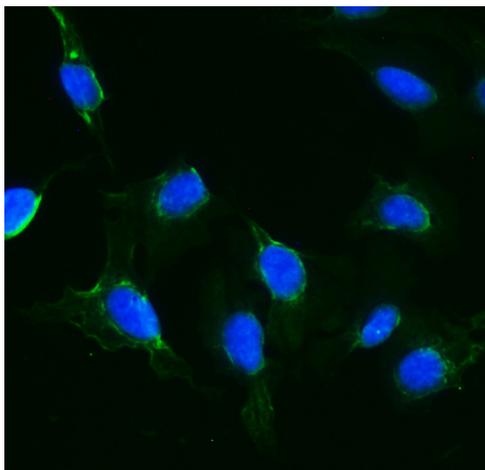
After electrophoresis, proteins were transferred to a membrane.

Then the membrane was incubated with rabbit anti-IFT88 antigen affinity purified polyclonal antibody (A06814-1) at a dilution of 1:1000 and probed with a goat anti-rabbit IgG-HRP secondary antibody (Catalog # BA1054). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for IFT88 at approximately 94 kDa. The expected band size for IFT88 is at 94 kDa.

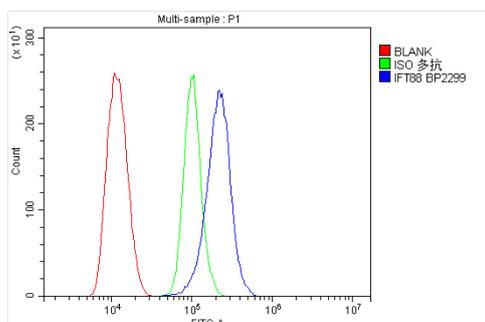


IHC analysis of IFT88 using anti-IFT88 antibody (A06814-1).

IFT88 was detected in a paraffin-embedded section of human renal cancer tissue. Biotinylated goat anti-rabbit IgG was used as secondary antibody. The tissue section was incubated with rabbit anti-IFT88 Antibody (A06814-1) at a dilution of 1:200 and developed using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB (Catalog # AR1027) as the chromogen.



ICC/IF analysis of IFT88 using anti-IFT88 antibody (A06814-1). IFT88 was detected in an immunocytochemical section of U2OS cells. The section was incubated with rabbit anti-IFT88 Antibody (A06814-1) at a dilution of 1:100. Fluoro488 Conjugated Goat Anti-Rabbit IgG (Green) (Catalog # BA1127) was used as secondary antibody. The section was counterstained with DAPI (Catalog # AR1176) (Blue).



Flow Cytometry analysis of THP-1 cells using anti-IFT88 antibody (A06814-1).

Overlay histogram showing THP-1 cells stained with A06814-1 (Blue line). To facilitate intracellular staining, cells were fixed with 4% paraformaldehyde and permeabilized with permeabilization buffer. The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-IFT88 Antibody (A06814-1) at 1:100 dilution for 30 min at 20°C. Fluoro488 conjugated goat anti-rabbit IgG (BA1127) was used as secondary antibody at 1:100 dilution for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG at 1:100 dilution used under the same conditions. Unlabelled sample without incubation with primary antibody and secondary antibody (Red line) was used as a blank control.