IFIT2 Antibody

PACO51630



Product Information	
Size:	Protein Background:
50ug	IFN-induced antiviral protein which inhibits expression of viral messenger RNAs lacking
Reactivity:	2'-O-methylation of the 5' cap. The ribose 2'-O-methylation would provide a molecular signature to distinguish between self and non-self mRNAs by the host during viral
Human, Rat	infection. Viruses evolved several ways to evade this restriction system such as encoding their own 2'-O-methylase for their mRNAs or by stealing host cap containing
Source:	the 2'-O-methylation (cap snatching mechanism). Binds AU-rich viral RNAs, with or
Rabbit	without 5' triphosphorylation, RNA-binding is required for antiviral activity. Can promote apoptosis.
lsotype:	Gene ID:
lgG	IFIT2
Applications:	Uniprot
ELISA, WB, IHC, IF	P09913
Recommended dilutions:	Synonyms:
ELISA:1:2000-1:10000, WB:1:500-1:5000, IHC:1:500-1:1000, IF:1:50-1:200	Interferon-induced protein with tetratricopeptide repeats 2 (IFIT-2) (ISG-54 K) (Interferon-induced 54 kDa protein) (IFI-54K) (P54), IFIT2, CIG-42 G10P2 IFI54 ISG54
	Immunogen:
	Recombinant Human Interferon-induced protein with tetratricopeptide repeats 2 protein (130-265AA).
	Storage:

Storage:

Preservative: 0.03% Proclin 300. Constituents: 50% Glycerol, 0.01M PBS, pH 7.4





Western Blot. Positive WB detected in: HEK293 whole cell lysate, Rat liver tissue. All lanes: IFIT2 antibody at 3µg/ml. Secondary. Goat polyclonal to rabbit IgG at 1/50000 dilution. Predicted band size: 55 kDa. Observed band size: 55 kDa.

IHC image of PACO51630 diluted at 1:600 and staining in paraffinembedded human colon cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.

IHC image of PACO51630 diluted at 1:600 and staining in paraffinembedded human cervical cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.