

## elF5B rabbit pAb

## Cat No.:ES8084

For research use only

## Overview

Product Name	eIF5B rabbit pAb
Host species	Rabbit
Applications	WB;IHC;IF;ELISA
Species Cross-Reactivity	Human;Mouse;Rat
Recommended dilutions	Western Blot: 1/500 - 1/2000.
	Immunohistochemistry: 1/100 - 1/300. ELISA:
	1/40000. Not yet tested in other applications.
Immunogen	The antiserum was produced against synthesized
	peptide derived from human EIF5B. AA
	range:1051-1100
Specificity	eIF5B Polyclonal Antibody detects endogenous
	levels of eIF5B protein.
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and
	0.02% sodium azide.
Storage	Store at -20°C. Avoid repeated freeze-thaw cycles.
Protein Name	Eukaryotic translation initiation factor 5B
Gene Name	EIF5B
Cellular localization	Cytoplasm .
Purification	The antibody was affinity-purified from rabbit
	antiserum by affinity-chromatography using
	epitope-specific immunogen.
Clonality	Polyclonal
Concentration	1 mg/ml
Observed band	139kD
Human Gene ID	9669
Human Swiss-Prot Number	O60841
Alternative Names	EIF5B; IF2; KIAA0741; Eukaryotic translation
	initiation factor 5B; eIF-5B; Translation initiation
	factor IF-2
Background	Accurate initiation of translation in eukaryotes is
	complex and requires many factors, some of which
	are composed of multiple subunits. The process is
	simpler in prokaryotes which have only three



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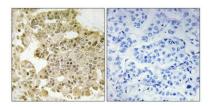
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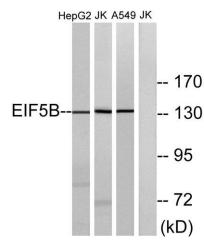
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initiation factors (IF1, IF2, IF3). Two of these factors are conserved in eukaryotes: the homolog of IF1 is eIF1A and the homolog of IF2 is eIF5B. This gene encodes eIF5B. Factors eIF1A and eIF5B interact on the ribosome along with other initiation factors and GTP to position the initiation methionine tRNA on the start codon of the mRNA so that translation initiates accurately. [provided by RefSeq, Jul 2008],

Immunohistochemical analysis of paraffin-embedded Human breast cancer. Antibody was diluted at 1:100(4° overnight). High-pressure and temperature Tris-EDTA,pH8.0 was used for antigen retrieval. Negetive contrl (right) obtaned from antibody was pre-absor





Western blot analysis of lysates from Jurkat, HepG2, and A549 cells, using EIF5B Antibody. The lane on the right is blocked with the synthesized peptide.



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