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# anti-PRPF31 antibody (Internal Region)

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**Images** 



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Overview	
Quantity:	100 μL
Target:	PRPF31
Binding Specificity:	Internal Region
Reactivity:	Human, Mouse, Rat
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This PRPF31 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Immunocytochemistry (ICC), Immunofluorescence (IF)

#### **Product Details**

Immunogen:	A synthesized peptide derived from human PRPF31, corresponding to a region within the internal amino acids.
Isotype:	IgG
Specificity:	PRPF31 Antibody detects endogenous levels of total PRPF31.
Predicted Reactivity:	Zebrafish,Bovine,Horse,Rabbit,Xenopus
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink <sup>TM</sup> Coupling Resin (Thermo Fisher Scientific).

### **Target Details**

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Target:	PRPF31	
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## **Target Details**

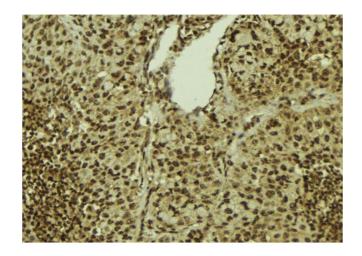
Alternative Name:	PRPF31 (PRPF31 Products)	
Background:	Description: Involved in pre-mRNA splicing as component of the spliceosome	
	(PubMed:11867543, PubMed:28781166). Required for the assembly of the U4/U5/U6 tri-snRNP	
	complex, one of the building blocks of the spliceosome (PubMed:11867543).	
	Gene: PRPF31	
Molecular Weight:	55kDa	
Gene ID:	26121	
UniProt:	Q8WWY3	
Pathways:	Ribonucleoprotein Complex Subunit Organization	

## **Application Details**

Application Notes:	WB 1:1000, IHC 1:200, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000	
Restrictions:	For Research Use only	

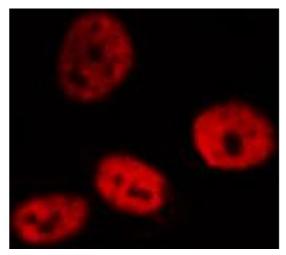
# Handling

Format:	Liquid
Concentration:	1 mg/mL
Buffer:	Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 % glycerol.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C
Storage Comment:	Store at -20 °C. Stable for 12 months from date of receipt.
Expiry Date:	12 months



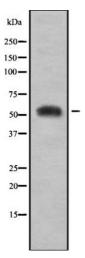
#### **Immunohistochemistry**

**Image 1.** ABIN6272401 at 1/100 staining Human breast cancer tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the antibody for 1.5 hours at 22°C. An HRP conjugated goat anti-rabbit antibody was used as the secondary.



#### Immunofluorescence (fixed cells)

**Image 2.** ABIN6272401 staining HeLa by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25°C. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary antibody.



#### **Western Blotting**

**Image 3.** Western blot analysis of PRPF31 using HeLa whole cell lysates