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anti-CSDC2 antibody (AA 76-105)

Datasheet for ABIN5532502





Overview	
Quantity:	400 μL
Target:	CSDC2
Binding Specificity:	AA 76-105
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This CSDC2 antibody is un-conjugated
Application:	Western Blotting (WB), Immunohistochemistry (Paraffin-embedded Sections) (IHC (p))
Product Details	
lmmunogen:	This PIPPIN antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 76-105 amino acids from the Central region of human PIPPIN.
Isotype:	Ig Fraction
Purification:	This antibody is purified through a protein A column, followed by peptide affinity purification.
Target Details	
Target:	CSDC2
Alternative Name:	PIPPIN (CSDC2 Products)
Background:	RNA-binding factor which binds specifically to the very 3'-UTR ends of both histone H1 and H3.3 mRNAs, encompassing the polyadenylation signal. Might play a central role in the negative regulation of histone variant synthesis in the developing brain (By similarity).

Target Details

Molecular Weight:	17 kDa
Gene ID:	27254
UniProt:	Q9Y534

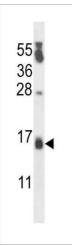
Application Details

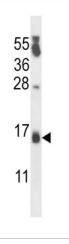
Application Notes:	For WB starting dilution is: 1:1000
	For IHC-P starting dilution is: 1:10~50
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Concentration:	0.5 mg/mL

Buffer:	Supplied in PBS with 0.09 % (W/V) sodium azide.
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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:	4 °C,-20 °C

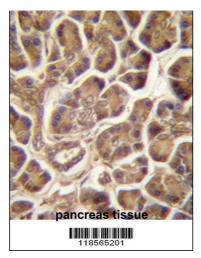
Storage Comment:	Store at 4°C for three months and -20°C, stable for up to one year. As with all antibodies care
	should be taken to avoid repeated freeze thaw cycles. Antibodies should not be exposed to
	prolonged high temperatures.





Western Blotting

Image 1. Western blot analysis in MDA-MB231 cell line lysates (35ug/lane).



Immunohistochemistry

Image 2. PIPPIN Antibody immunohistochemistry analysis in formalin fixed and paraffin embedded human pancreas tissue followed by peroxidase conjugation of the secondary antibody and DAB staining.