

**Anti-SNRPN Picoband Antibody**  
Catalog # ABO12131**Specification****Anti-SNRPN Picoband Antibody - Product Information**

Application	WB, IHC
Primary Accession	<a href="#">P63162</a>
Host	Rabbit
Reactivity	Human, Mouse, Rat
Clonality	Polyclonal
Format	Lyophilized

**Description**

Rabbit IgG polyclonal antibody for Small nuclear ribonucleoprotein-associated protein N(SNRPN) detection. Tested with WB, IHC-P in Human;Mouse;Rat.

**Reconstitution**

Add 0.2ml of distilled water will yield a concentration of 500ug/ml.

**Anti-SNRPN Picoband Antibody - Additional Information**

**Gene ID** 6638;8926

**Other Names**

Small nuclear ribonucleoprotein-associated protein N, snRNP-N, Sm protein D, Sm-D, Sm protein N, Sm-N, SmN, Tissue-specific-splicing protein, SNRPN, HCERN3, SMN

**Calculated MW**

24614 MW KDa

**Application Details**

Immunohistochemistry(Paraffin-embedded Section), 0.5-1 µg/ml, Human, Mouse, Rat, By Heat<br>Western blot, 0.1-0.5 µg/ml, Human, Mouse, Rat<br>

**Subcellular Localization**

Nucleus.

**Tissue Specificity**

Expressed in brain and lymphoblasts. .

**Protein Name**

Small nuclear ribonucleoprotein-associated protein N

**Contents**

Each vial contains 5mg BSA, 0.9mg NaCl, 0.2mg Na<sub>2</sub>HPO<sub>4</sub>, 0.05mg Na<sub>3</sub>.

**Immunogen**

A synthetic peptide corresponding to a sequence at the N-terminus of human SNRPN (11-39aa QHIDYRMRCILQDGRIFIGTFKAFDKHMN), identical to the related mouse and rat sequences.

**Purification**

Immunogen affinity purified.

**Cross Reactivity**

No cross reactivity with other proteins

**Storage**

At -20°C for one year. After reconstitution, at 4°C for one month. It can also be aliquotted and stored frozen at -20°C for a longer time. Avoid repeated freezing and thawing.

**Anti-SNRPN Picoband Antibody - Protein Information**

**Name** SNRPN

**Synonyms** HCERN3, SMN

**Function**

May be involved in tissue-specific alternative RNA processing events.

**Cellular Location**

Nucleus.

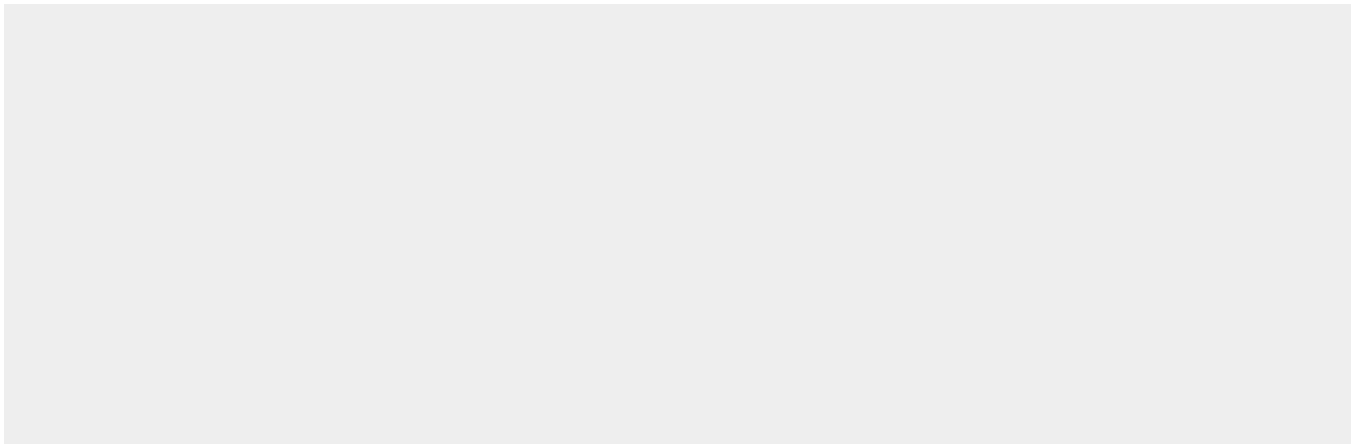
**Tissue Location**

Expressed in brain and lymphoblasts.

**Anti-SNRPN Picoband Antibody - Protocols**

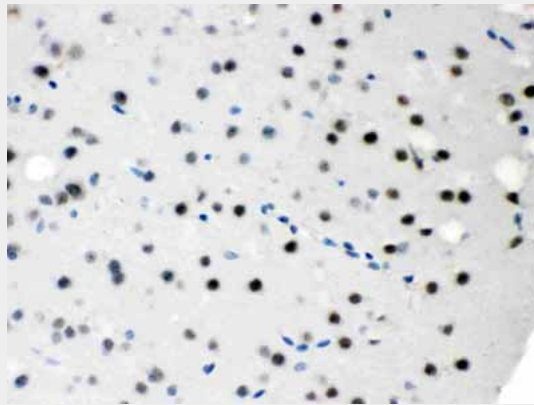
Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

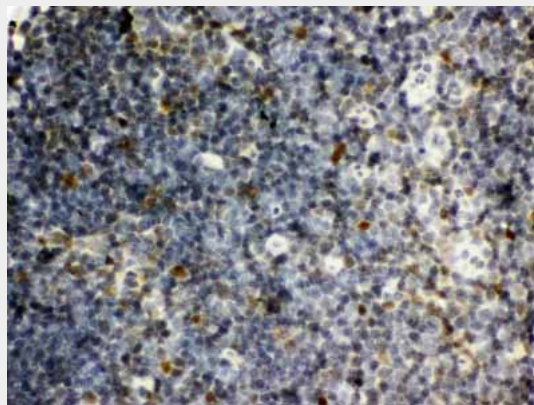
**Anti-SNRPN Picoband Antibody - Images**



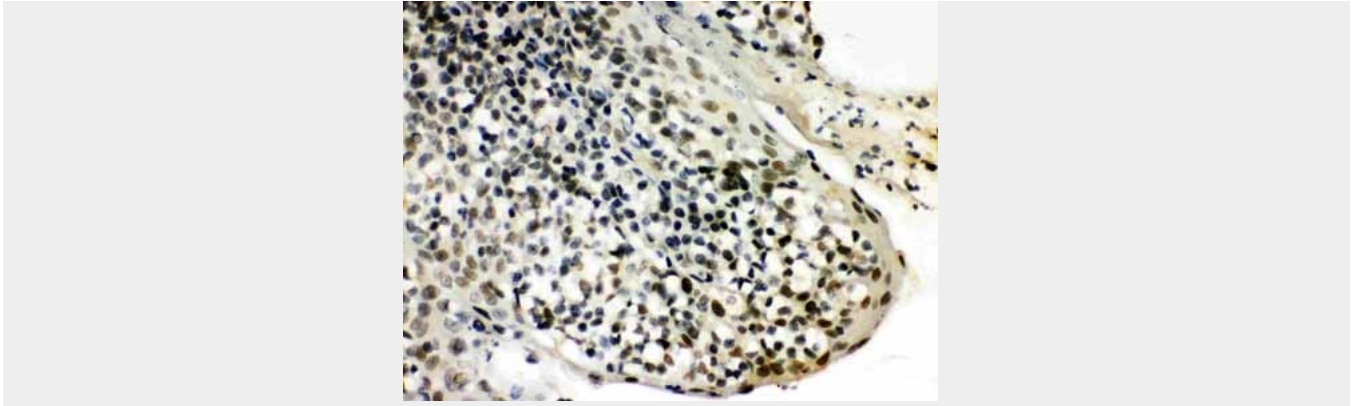
Anti- SNRPN Picoband antibody, ABO12131, Western blotting All lanes: Anti SNRPN (ABO12131) at 0.5ug/ml Lane 1: Rat Thymus Tissue Lysate at 50ug Lane 2: HELA Whole Cell Lysate at 40ug Lane 3: HEPG2 Whole Cell Lysate at 40ug Lane 4: HEPA Whole Cell Lysate at 40ug Predicted bind size: 25KD Observed bind size: 32KD



Anti- SNRPN Picoband antibody, ABO12131, IHC(P) IHC(P): Mouse Brain Tissue



Anti- SNRPN Picoband antibody, ABO12131, IHC(P) IHC(P): Rat Thymus Tissue



Anti- SNRPN Picoband antibody, ABO12131, IHC(P)IHC(P): Human Tonsil Tissue

### **Anti-SNRPN Picoband Antibody - Background**

SNRPN (Small Nuclear Ribonucleoprotein Polypeptide N), also called SMN, is a bicistronic imprinted gene that encodes 2 polypeptides, the SmN splicing factor, which is involved in RNA processing, and the SNRPN upstream reading frame (SNURF) polypeptide. The protein encoded by this gene is one polypeptide of a small nuclear ribonucleoprotein complex and belongs to the snRNP SMB/SMN family. SNRPN also encodes a long alternatively spliced transcript containing several small nucleolar RNAs (snoRNAs) and extends downstream to partially overlap the UBE3A gene in the antisense orientation. PWS arises from loss of function of genes in this region expressed exclusively from the paternal chromosome, suggesting that SNRPN may play a role in its etiology. The SNRPN gene is mapped on 15q11.2. Analysis of maternal DNA and of SNRPN cDNA confirmed that the maternal allele is not expressed in fetal brain and heart. Deletions in the transcription unit of the imprinted SNRPN gene occur in patients who have PWS or Angelman syndrome because of a parental imprint switch failure in this chromosomal domain.