

CAB15726

Product Information

| Product SKU: | CAB15726 | Gene ID: | 6638 | | Size : | 20uL, 100uL | | |
|------------------------|----------|---------------|------------|--------------|---------------------|-------------------|--|--|
| Clone No: | - | Host Species: | Rabbit | | Reactivity : | Human, Mouse, Rat | | |
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| Additional Information | | | | | | | | |
| Observed MW: | 20kDa | | Conjugate: | Unconjugated | Ł | | | |

| Calculated MW: | 25kDa | lsotype: | lgG |
|----------------|-------|----------|-----|
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Immunogen Information

| Background: This gene is located within the Prader-Willi Syndrome critical region on chromosome 15 and is imprinted and expressed from the paternal allele. It encodes a component of the small nuclear ribonucleoprotein complex, which functions in pre-mRNA processing and may contribute to tissue-specific alternative |
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| complex, which functions in pre-mRNA processing and may contribute to tissue-specific alternative |
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| splicing. Alternative promoter use and alternative splicing result in a multitude of transcript variants |
| encoding the same protein. Transcript variants that initiate at the CpG island-associated imprinting |
| center may be bicistronic and also encode the SNRPN upstream reading frame protein (SNURF) from an |
| upstream open reading frame. In addition, long spliced transcripts for small nucleolar RNA host gene |
| 14 (SNHG14) may originate from the promoters at this locus and share exons with this gene. Alterations |
| in this region are associated with parental imprint switch failure, which may cause Angelman syndrome |
| or Prader-Willi syndrome. |
| Recommended Dilution: WB,1:500 - 1:2000 |
| Synonyms: SMN; PWCR; SM-D; sm-N; RT-LI; HCERN3; SNRNP-N; SNURF-SNRPN; SNRPN |
| Purifcation Method: Affinity purification |
| Immunogen: A synthetic peptide corresponding to a sequence within amino acids 1-100 of human SNRPN |
| (NP_003088.1). |
| Storage: Store at -20°C. Avoid freeze / thaw cycles.Buffer: PBS with 0.01% thimerosal,50% glycerol,pH7.3. |