Product Sheet

• pN0.9 95679™

Description

Detects: Genome: Homo sapiens Gene symbol: SNRPN Type of nucleic acid: genomic Restriction enzyme: Xbal/Notl Number of alleles: 2 Type of polymorphism: site Alleles: Allele Freq. Size (kb) Strains 4.2 Xbal fragment with 3 internal methylated Notl sites 0.9 Notl subfragment resulting from absence of methylation at 3 Notl sites **Organism:** Homo sapiens, human Clone type: Clone Shipping information: Rehydrate with TE

Storage Conditions Product format: Dried

Intended Use

This product is intended for laboratory research use only. It is not intended for any animal or human therapeutic use, any human or animal consumption, or any diagnostic use.



BSL 1

ATCC determines the biosafety level of a material based on our risk assessment as guided by the current edition of *Biosafety in Microbiological and Biomedical Laboratories (BMBL)*, U.S. Department of Health and Human Services. It is your responsibility to understand the hazards associated with the material per your organization's policies and procedures as well as any other applicable regulations as enforced by your local or national agencies.

Certificate of Analysis

For batch-specific test results, refer to the applicable certificate of analysis that can be found at www.atcc.org.

Insert Information

Insert size (kb): 0.9000000000000002 Type of DNA: genomic Insert information: **DESCRIPTION OF INSERT COMPONENT:** Cross references: DNA Seq. Acc.: L32702 Nucleotides 1-196 of the insert correspond to nucleotides 216-411 of L32702. Genome: Homo sapiens Chromosome: 15 15 q11-q12 Target gene: Prader-Willi syndrome chromosome region, small nuclear ribonucleoprotein polypeptide N Gene name: small nuclear ribonucleoprotein polypeptide N Gene product: small nuclear ribonucleoprotein polypeptide N(small nuclear ribonucleoprotein SM-D, small nuclear ribonucleoprotein polypeptide Sm-D) [SNRPN] Gene symbol: SNRPN Contains complete coding sequence: No



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Insert end: Notl

Vector Information

Construct size (kb): 3.90000095367432 Intact vector size: 2.961 Vector name: pBluescript II SK-Type of vector: phagemid Host range: Escherichia coli Vector end: Notl Cloning sites: BssHII; KpnI; Apal; Drall; XhoI; HincII; AccI; SalI; ClaI; HindIII; EcoRI; PstI; Smal; BamHI; XbaI; NotI; EagI; SacI Insert detection: lacZ' Markers: ampR Polylinker sites: BssHII; KpnI; ApaI; DralI; XhoI; HincII; AccI; SalI; ClaI; HindIII; EcoRI; PstI; Smal; BamHI; XbaI; NotI; EagI; SacI Insert detection: lacZ' Markers: ampR

Notes

Restriction digests of the clone give the following sizes (kb): BamHI--4.0; EcoRI--3.6, 0.33; NotI--2.9, 0.88; PvuII--2.6, 1.0 0.33; XhoI--4.0. - ATCC staff

Insert contains a deletion breakpoint found in some cases of familial Prader-Willi syndrome (PWS). - Nat. Genet. 8: 52-58, 1994

The insert contains most of exon alpha (nt 1-66), also corresponding to nt 10-75 of GenBank accession J04615, and a portion of the following intron. - Nat. Genet. 8: 52-58, 1994

The 0.9 kb Notl fragment is derived from a 4.2 kb Xbal genomic DNA fragment

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which shows differential methylation at several rare restriction sites in normal maternal and paternal chromosomes. - Nat. Genet. 8: 52-58, 1994

Insert can be used to probe genomic DNA digested with Notl/Xbal to detect abnormalities diagnostic for Prader-Willi syndrome and Angelman syndrome (AS). - Nat. Genet. 8: 52-58, 1994

The probe detects a 4.2 kb fragment for normal maternal chromosomes, consistent with methylation of three NotI sites. An 0.9 kb fragment is detected for normal paternal chromosomes, consistent with the absence of methylation at these NotI sites.

- Nat. Genet. 8: 52-58, 1994

Normal individuals will show fragments of both sizes. Absence of the 0.9 kb band is indicative of PWS. Absence of the 4.2 kb band is indicative of AS. - Nat. Genet. 8: 52-58, 1994

Insert contains the following restriction sites (approximate kb from the 5' end): EcoRI--0.17, 0.63; PuvII--0.07. - Nat. Genet. 8: 52-58, 1994

Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: pN0.9 (ATCC 95679)

References

References and other information relating to this material are available at

www.atcc.org.

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