

pPW71B [PW71B]

99413™

Description

Detects:

Genome: Homo sapiens Gene symbol: D15S63

Type of nucleic acid: dsDNA

Restriction enzyme: HindIII+Hpall

Number of alleles: 2

Type of polymorphism: parental methylation

Alleles:

Allele Freq. Size (kb) Strains 4.7 paternal chromosomes 6.6 maternal chromosomes

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.

BSL₁



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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): 0.36499999999999999

Type of DNA: genomic **Genome:** Homo sapiens

Chromosome: 15

15 q11-q12

Target gene: DNA Segment, single copy probes **Gene name:** DNA Segment, single copy probes

Gene product: DNA Segment, single copy probes [D15S63]

Gene symbol: D15S63

Contains complete coding sequence: No

Insert end: Haelll

Vector Information

Construct size (kb): 3.049999952316284

Intact vector size: 2.686 Vector name: pUC19 Type of vector: plasmid



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Construction: pUC71K
Host range: Escherichia coli

Vector end: Smal Vector information:

Cross references: DNA Seq. Acc.: X02514

Cloning sites: EcoRI; SacI; KpnI; SmaI; BamHI; XbaI; HincII; AccI; SalI; PstI; SphI; HindIII

Insert detection: lacZ', <-</pre>

Markers: ampR

MCS: EcoRI...HindIII, ->

Polylinker sites: EcoRl; Sacl; Kpnl; Smal; BamHl; Xbal; Hincll; Accl; Sall; Pstl; Sphl; Hindlll

Replicon: pMB1, <-

Notes

Restriction digests of the clone give the following sizes (kb): EcoRI/HindIII--2.8, 0.44; PstI--3.2, PvuII--2.5, 0.76; SacI--3.2; XbaI--3.2.

- ATCC staff

Insert detects parent-of-origin specific DNA methylation patterns in the Prader-Willi syndrome/Angelman syndrome chromosomal region (15 q11-q13). An imprinted Hpall and Cfol site in this region results in the maternal and paternal pattern.

- Hum. Mol. Genet. 2: 1995-1999, 1993

Patients with Prader-Willi syndrome typically lack the paternal bands. Lack of the maternal bands is indicative of Angelman syndrome. Because of the small probe size, low-stringent post-hybridization washes should be applied.

- Hum. Mol. Genet. 2: 1995-1999, 1993

Material Citation

If use of this material results in a scientific publication, please cite the material in the

following manner: pPW71B [PW71B] (ATCC 99413)

References

References and other information relating to this material are available at www.atcc.org.

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