C19orf47 (S-22): sc-130990



The Power to Question

BACKGROUND

Consisting of around 63 million bases with over 1,400 genes, chromosome 19 makes up over 2% of human genomic DNA. Chromosome 19 includes a diversity of interesting genes and is recognized for having the greatest gene density of the human chromosomes. It is the genetic home for a number of immunoglobulin superfamily members including the killer cell and leukocyte lg-like receptors, a number of ICAMs, the CEACAM and PSG family, and Fc α receptors. Key genes for eye color and hair color also map to chromosome 19. Peutz-Jeghers syndrome, spinocerebellar ataxia type 6, the stroke disorder CADASIL, hypercholesterolemia and Insulin-dependent diabetes have been linked to chromosome 19. Translocations with chromosome 19 and chromosome 14 can be seen in some lymphoproliferative disorders and typically involve the proto-oncogene Bcl-3. The C19orf47 gene product has been provisionally designated C19orf47 pending further characterization.

REFERENCES

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- 3. Trettel, F., et al. 2000. A fine physical map of the CACNA1A gene region on 19p13.1-p13.2 chromosome. Gene 241: 45-50.
- 4. Buchet-Poyau, K., et al. 2002. Search for the second Peutz-Jeghers syndrome locus: exclusion of the STK13, PRKCG, KLK10, and PSCD2 genes on chromosome 19 and the STK11IP gene on chromosome 2. Cytogenet. Genome Res. 97: 171-178.
- 5. Moodie, S.J., et al. 2002. Analysis of candidate genes on chromosome 19 in coeliac disease: an association study of the KIR and LILR gene clusters. Eur. J. Immunogenet. 29: 287-291.
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CHROMOSOMAL LOCATION

Genetic locus: C19orf47 (human) mapping to 19q13.2; 2310022A10Rik (mouse) mapping to 7 A3.

SOURCE

C19orf47 (S-22) is an affinity purified rabbit polyclonal antibody raised against synthetic C19orf47 peptide of human origin.

PRODUCT

Each vial contains 50 μg lgG in 500 μl PBS with < 0.1% sodium azide, 0.1% gelatin and < 0.02% sucrose.

RESEARCH USE

For research use only, not for use in diagnostic procedures.

APPLICATIONS

C19orf47 (S-22) is recommended for detection of C19orf47 of human origin and 2310022A10Rik of mouse origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)] and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for C19orf47 siRNA (h): sc-97383, 2310022A10Rik siRNA (m): sc-108674, C19orf47 shRNA Plasmid (h): sc-97383-SH, 2310022A10Rik shRNA Plasmid (m): sc-108674-SH, C19orf47 shRNA (h) Lentiviral Particles: sc-97383-V and 2310022A10Rik shRNA (m) Lentiviral Particles: sc-108674-V.

Molecular Weight (predicted) of C19orf47: 45 kDa.

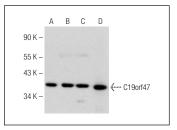
Molecular Weight (observed) of C19orf47: 37 kDa.

Positive Controls: Jurkat whole cell lysate: sc-2204, LNCaP cell lysate: sc-2231 or NAMALWA cell lysate: sc-2234.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western blotting: use goat anti-rabbit IgG-HRP: sc-2004 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible goat anti-rabbit IgG-HRP: sc-2030 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml).

DATA



C19orf47 (S-22): sc-130990. Western blot analysis of C19orf47 expression in LNCaP (A), NAMALWA (B) and Jurkat (C) whole cell lysates and mouse skeletal muscle tissue extract (D).

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.



Try **C19orf47 (H-5): sc-393896**, our highly recommended monoclonal alternative to C19orf47 (S-22).