C21orf56 (P-17): sc-83549



The Power to Question

BACKGROUND

The smallest of the human chromosomes, 21 makes up about 1.5% of the human genome. Chromosome 21 contains nearly 300 genes and 47 million base pairs. Down syndrome, also known as trisomy 21, is the disease most commonly associated with chromosome 21. Alzheimer's disease, Jervell and Lange-Nielsen syndrome, and amyotrophic lateral sclerosis are also associated with chromosome 21. Translocations are found to occur between chromosome 21 and 8, and chromosome 21 and 12 in certain leukemias. The C21orf56 gene product has been provisionally designated C21orf56 pending further characterization.

REFERENCES

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- Mao, R., et al. 2005. Primary and secondary transcriptional effects in the developing human Down syndrome brain and heart. Genome Biol. 6: R107.
- Robakis, N.K. 2006. The discovery and mapping to chromosome 21 of the Alzheimer's amyloid gene: history revised. J. Alzheimers Dis. 10: 453-455.
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CHROMOSOMAL LOCATION

Genetic locus: C21orf56 (human) mapping to 21q22.3; 1700027D21Rik (mouse) mapping to 10 C1.

SOURCE

C21orf56 (P-17) is an affinity purified rabbit polyclonal antibody raised against a peptide mapping within an internal region of C21orf56 of human origin.

PRODUCT

Each vial contains 100 μg lgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-83549 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

STORAGE

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

APPLICATIONS

C21orf56 (P-17) is recommended for detection of C21orf56 of human and mouse origin and the corresponding rat homolog by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

C21orf56 (P-17) is also recommended for detection of C21orf56 in additional species, including equine, bovine and porcine.

Suitable for use as control antibody for C21orf56 siRNA (h): sc-91465, C21orf56 siRNA (m): sc-108400, C21orf56 shRNA Plasmid (h): sc-91465-SH, C21orf56 shRNA Plasmid (m): sc-108400-SH, C21orf56 shRNA (h) Lentiviral Particles: sc-91465-V and C21orf56 shRNA (m) Lentiviral Particles: sc-108400-V.

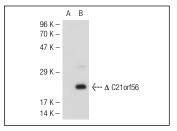
Molecular Weight of C21orf56: 38 kDa.

Positive Controls: C21orf56 (h): 293T Lysate: sc-112414.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use goat anti-rabbit lgG-HRP: sc-2004 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible goat anti-rabbit lgG-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). 3) Immunofluorescence: use goat anti-rabbit lgG-FITC: sc-2012 (dilution range: 1:100-1:400) or goat anti-rabbit lgG-TR: sc-2780 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

DATA



C21orf56 (P-17): sc-83549. Western blot analysis of truncated C21orf56 expression in non-transfected: sc-117752 (A) and human C21orf56 transfected: sc-112414 (B) 2937 whole cell lysates.

RESEARCH USE

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

PROTOCOLS

See our web site at www.scbt.com or our catalog for detailed protocols and support products.