LRRC8A (8H9): sc-517113



The Power to Question

BACKGROUND

Chromosome 9 consists of about 145 million bases and 4% of the human genome, encoding nearly 900 genes. Considered to play a role in gender determination, deletion of the distal portion of 9p can lead to development of male to female sex reversal, the phenotype of a female with a male X,Y genotype. Hereditary hemorrhagic telangiectasia, which is characterized by harmful vascular defects, is associated with the chromosome 9 gene-encoding Endoglin protein, ENG. Familial dysautonomia is also associated with chromosome 9 though through the gene IKBKAP. Notably, chromosome 9 encompasses the largest interferon family gene cluster. Chromosome 9 is partnered with chromosome 22 in the translocation leading to the aberrant production of Bcr-Abl fusion protein often found in leukemias.

REFERENCES

- Fryns, J.P., et al. 1991. Apparent late-onset Cockayne syndrome and interstitial deletion of the long arm of chromosome 10 (del(10)(q11.23q21.2)).
 Am. J. Med. Genet. 40: 343-344.
- 2. Thöny, B., et al. 1994. Chromosomal location of two human genes encoding tetrahydrobiopterin-metabolizing enzymes: 6-pyruvoyl-tetrahydropterin synthase maps to 11q22.3-q23.3, and pterin-4 α -carbinolamine dehydratase maps to 10q22. Genomics 19: 365-368.

CHROMOSOMAL LOCATION

Genetic locus: LRRC8A (human) mapping to 9q34.11; Lrrc8a (mouse) mapping to 2 B.

SOURCE

LRRC8A (8H9) is a mouse monoclonal antibody raised against amino acids 711-810 representing partial length LRRC8A of human origin.

PRODUCT

Each vial contains 100 μg lgG_{2a} kappa light chain in 1.0 ml PBS with < 0.1% sodium azide and 0.1% gelatin.

STORAGE

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

APPLICATIONS

LRRC8A (8H9) is recommended for detection of LRRC8A of mouse, rat and human origin 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)] and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

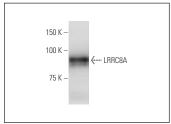
Suitable for use as control antibody for LRRC8A siRNA (h): sc-92566, LRRC8A siRNA (m): sc-149105, LRRC8A shRNA Plasmid (h): sc-92566-SH, LRRC8A shRNA Plasmid (m): sc-149105-SH, LRRC8A shRNA (h) Lentiviral Particles: sc-92566-V and LRRC8A shRNA (m) Lentiviral Particles: sc-149105-V.

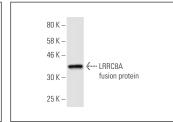
Positive Controls: A-431 whole cell lysate: sc-2201.

RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-lgG κ BP-HRP: sc-516102 or m-lgG κ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz MarkerTM Molecular Weight Standards: sc-2035, UltraCruz® Blocking Reagent: sc-516214 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





LRRC8A (8H9); sc-517113. Western blot analysis of LRRC8A expression in A-431 whole cell lysate.

LRRC8A (8H9): sc-517113. Western blot analysis of human recombinant LRRC8A fusion protein.

SELECT PRODUCT CITATIONS

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- 2. Zhang, H., et al. 2019. Novel insights into the role of LRRC8A in ameliorating alveolar fluid clearance in LPS induced acute lung injury. Eur. J. Pharmacol. 861: 172613.
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- Cook, J.R., et al. 2022. LRRC8A is dispensable for a variety of microglial functions and response to acute stroke. Glia 70: 1068-1083.
- 5. Liu, J., et al. 2022. Inhibition of the LRRC8A channel promotes microglia/macrophage phagocytosis and improves outcomes after intracerebral hemorrhagic stroke. iScience 25: 105527.
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- 7. Liu, Y., et al. 2023. Interactions between the astrocytic volume-regulated anion channel and aquaporin 4 in hyposmotic regulation of vasopressin neuronal activity in the supraoptic nucleus. Cells 12: 1723.
- Balkaya, M., et al. 2023. Conditional deletion of LRRC8A in the brain reduces stroke damage independently of swelling-activated glutamate release. iScience 26: 106669.

RESEARCH USE

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