CLIP-115 siRNA (m): sc-60476



The Power to Question

BACKGROUND

Williams syndrome (WS) is a developmental disorder characterized by cardiovascular problems, dysmorphic features, mental retardation or learning difficulties and several typical behavioral and neurological abnormalities. In Williams syndrome patients, a heterozygous deletion is present in a region on chromosome 7q11.23 (the Williams syndrome critical region), which spans approximately 20 genes. This region comprises the CYLN2 gene, which encodes the cytoplasmic linker protein of (CLIP-115). CLIP-115 is a microtubule-binding protein that is abundantly expressed in the brain. Mice with haploin-sufficiency for the CYLN2 gene have features similar to that of WS, including mild growth deficiency, brain abnormalities, hippocampal dysfunction and particular deficits in motor coordination.

REFERENCES

- 1. Hoogenraad, C.C., et al. 1998. The murine CYLN2 gene: genomic organization, chromosome localization, and comparison to the human gene that is located within the 7q11.23 Williams syndrome critical region. Genomics 53: 348-358.
- 2. Donnai, D., et al. 2001. Williams syndrome: from genotype through to the cognitive phenotype. Am. J. Med. Genet. 97: 164-171.
- 3. Hoogenraad, C.C., et al. 2002. Targeted mutation of Cyln2 in the Williams syndrome critical region links CLIP-115 haploinsufficiency to neurodevelopmental abnormalities in mice. Nat. Genet. 32: 116-127.
- Galaburda, A.M., et al. 2003. Williams syndrome. A summary of cognitive, electrophysiological, anat microanatomical and genetic findings. Rev. Neurol. 1: S132-S137.
- 5. Hoogenraad, C.C., et al. 2004. LIMK1 and CLIP-115: linking cytoskeletal defects to Williams syndrome. Bioessays 26: 141-150.
- Meyer-Lindenberg, A., et al. 2005. Functional, structural, and metabolic abnormalities of the hippocampal formation in Williams syndrome. J. Clin. Invest. 115: 1888-1895.
- 7. Meyer-Lindenberg, A., et al. 2006. Neural mechanisms in Williams syndrome: a unique window to genetic influences on cognition and behaviour. Nat. Rev. Neurosci. 7: 380-393.

CHROMOSOMAL LOCATION

Genetic locus: Clip2 (mouse) mapping to 5 G2.

PRODUCT

CLIP-115 siRNA (m) is a pool of 3 target-specific 19-25 nt siRNAs designed to knock down gene expression. Each vial contains 3.3 nmol of lyophilized siRNA, sufficient for a 10 μM solution once resuspended using protocol below. Suitable for 50-100 transfections. Also see CLIP-115 shRNA Plasmid (m): sc-60476-SH and CLIP-115 shRNA (m) Lentiviral Particles: sc-60476-V as alternate gene silencing products.

For independent verification of CLIP-115 (m) gene silencing results, we also provide the individual siRNA duplex components. Each is available as 3.3 nmol of lyophilized siRNA. These include: sc-60476A, sc-60476B and sc-60476C.

STORAGE AND RESUSPENSION

Store lyophilized siRNA duplex at -20 $^{\circ}$ C with desiccant. Stable for at least one year from the date of shipment. Once resuspended, store at -20 $^{\circ}$ C, avoid contact with RNAses and repeated freeze thaw cycles.

Resuspend lyophilized siRNA duplex in 330 μ l of the RNAse-free water provided. Resuspension of the siRNA duplex in 330 μ l of RNAse-free water makes a 10 μ M solution in a 10 μ M Tris-HCl, pH 8.0, 20 mM NaCl, 1 mM EDTA buffered solution.

APPLICATIONS

CLIP-115 siRNA (m) is recommended for the inhibition of CLIP-115 expression in mouse cells.

SUPPORT REAGENTS

For optimal siRNA transfection efficiency, Santa Cruz Biotechnology's siRNA Transfection Reagent: sc-29528 (0.3 ml), siRNA Transfection Medium: sc-36868 (20 ml) and siRNA Dilution Buffer: sc-29527 (1.5 ml) are recommended. Control siRNAs or Fluorescein Conjugated Control siRNAs are available as 10 µM in 66 µl. Each contain a scrambled sequence that will not lead to the specific degradation of any known cellular mRNA. Fluorescein Conjugated Control siRNAs include: sc-36869, sc-44239, sc-44240 and sc-44241. Control siRNAs include: sc-37007, sc-44230, sc-44231, sc-44232, sc-44233, sc-44234, sc-44235, sc-44236, sc-44237 and sc-44238.

GENE EXPRESSION MONITORING

CLIP-115 (14): sc-135869 is recommended as a control antibody for monitoring of CLIP-115 gene expression knockdown by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000) or immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500).

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) Immunofluorescence: use m-lgG κ BP-FITC: sc-516140 or m-lgG κ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz[®] Mounting Medium: sc-24941 or UltraCruz[®] Hard-set Mounting Medium: sc-359850.

RT-PCR REAGENTS

Semi-quantitative RT-PCR may be performed to monitor CLIP-115 gene expression knockdown using RT-PCR Primer: CLIP-115 (m)-PR: sc-60476-PR (20 μ l). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.

RESEARCH USE

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

PROTOCOLS

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

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