TEX11 (E-12): sc-240991



The Power to Question

BACKGROUND

TEX11 (testis-expressed sequence 11 protein) is a 940 amino acid protein that exists as 3 alternatively spliced isoforms. Forming discrete foci on synapsed regions of meiotic chromosomes, TEX11 may be a constituent of meiotic nodules involved in recombination. TEX11 is thought to promote initiation and/or maintenance of synapsis and formation of crossovers, and may provide a link between these two meiotic processes. The gene the encodes TEX11 consists of nearly 380,000 bases and maps to human chromosome Xq13.1. Chromosome X consists of about 153 million base pairs and nearly 1,000 genes. There are a number of conditions related to an unusual number and combination of sex chromosomes being inherited, including Turner's syndrome, Klinefelter's syndrome and Triple X syndrome. Color blindness, hemophilia, and Duchenne muscular dystrophy are well known X chromosome-linked conditions which affect males more frequently as males carry a single X chromosome.

REFERENCES

- Online Mendelian Inheritance in Man, OMIM™. 2001. Johns Hopkins University, Baltimore, MD. MIM Number: 300311. World Wide Web URL: http://www.ncbi.nlm.nih.gov/omim/
- Bernardino-Sgherri, J., et al. 2002. Overall DNA methylation and chromatin structure of normal and abnormal X chromosomes. Cytogenet. Genome Res. 99: 85-91.
- 3. Deeb, S.S. 2005. The molecular basis of variation in human color vision. Clin. Genet. 67: 369-377.
- 4. Bojesen, A., et al. 2006. The metabolic syndrome is frequent in Klinefelter's syndrome and is associated with abdominal obesity and hypogonadism. Diabetes Care 29: 1591-1598.
- Maggio, M.C., et al. 2007. Polycystic ovary and gonadoblastoma in Turner's syndrome. Minerva Pediatr. 59: 397-401.
- Yang, F., et al. 2008. Meiotic failure in male mice lacking an X-linked factor. Genes Dev. 22: 682-691.
- Helderman-van den Enden, A.T., et al. 2009. Recurrence risk due to germ line mosaicism: Duchenne and Becker muscular dystrophy. Clin. Genet. 75: 465-472.
- Kasper, C.K., et al. 2009. Mosaicism and haemophilia. Haemophilia. E-published.

CHROMOSOMAL LOCATION

Genetic locus: Tex11 (mouse) mapping to X C3.

SOURCE

TEX11 (E-12) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the N-terminus of TEX11 of mouse origin.

STORAGE

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

PRODUCT

Each vial contains 200 μg lgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-240991 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

TEX11 (E-12) is recommended for detection of TEX11 of mouse and rat origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), 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); non cross-reactive with other TEX family members.

Suitable for use as control antibody for TEX11 siRNA (m): sc-154210, TEX11 shRNA Plasmid (m): sc-154210-SH and TEX11 shRNA (m) Lentiviral Particles: sc-154210-V.

Molecular Weight of TEX11 isoforms: 108/71/107 kDa.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (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) Immunofluorescence: use donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

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.

Santa Cruz Biotechnology, Inc. 1.800.457.3801 831.457.3801 **Fax** 831.457.3801 **Europe** +00800 4573 8000 49 6221 4503 0 **www.scbt.com**