

# **SMCP Antibody**

Purified Mouse Monoclonal Antibody Catalog # AO1905a

### **Product Information**

**Application** WB, E **Primary Accession** P49901 Reactivity Human Host Mouse Monoclonal Clonality **Clone Names** 5C10A2 Isotype IgG2b 12767 **Calculated MW** 

**Description** Sperm mitochondria differ in morphology and subcellular localization from

those of somatic cells. They are elongated, flattened, and arranged

circumferentially to form a helical coiled sheath in the midpiece of the sperm

flagellum. The protein encoded by this gene localizes to the capsule associated with the mitochondrial outer membranes and is thought to function in the organization and stabilization of the helical structure of the

sperm's mitochondrial sheath.

**Immunogen** Purified recombinant fragment of human SMCP (AA: 1-116) expressed in E.

Coli.

**Formulation** Purified antibody in PBS with 0.05% sodium azide.

## **Additional Information**

**Gene ID** 4184

Other Names Sperm mitochondrial-associated cysteine-rich protein, SMCP, MCS, MCSP

**Dilution** WB~~1/500 - 1/2000 E~~1/10000

**Storage** Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store

at -20°C in small aliquots to prevent freeze-thaw cycles.

**Precautions** SMCP Antibody is for research use only and not for use in diagnostic or

therapeutic procedures.

#### **Protein Information**

Name SMCP

**Synonyms** MCS, MCSP

**Function** Involved in sperm motility. Its absence is associated with genetic

background dependent male infertility. Infertility may be due to reduced sperm motility in the female reproductive tract and inability to penetrate the

oocyte zona pellucida (By similarity).

**Cellular Location** Cytoplasm. Mitochondrion membrane; Peripheral membrane protein;

Cytoplasmic side. Note=Becomes associated with the spermatid

mitochondrion capsule at step 16 of spermatogenesis.

**Tissue Location** Testis. Is selectively expressed in the spermatids of seminiferous tubules.

# **Background**

Mutations in this gene are associated with Nijmegen breakage syndrome, an autosomal recessive chromosomal instability syndrome characterized by microcephaly, growth retardation, immunodeficiency, and cancer predisposition. The encoded protein is a member of the MRE11/RAD50 double-strand break repair complex which consists of 5 proteins. This gene product is thought to be involved in DNA double-strand break repair and DNA damage-induced checkpoint activation.;

#### References

1. Biol Reprod. 1999 Aug;61(2):428-35. 2. Mol Cell Biol. 2002 May;22(9):3046-52.

## **Images**

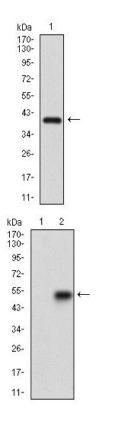


Figure 1: Western blot analysis using SMCP mAb against human SMCP (AA: 1-116) recombinant protein. (Expected MW is 38.3 kDa)

Figure 2: Western blot analysis using SMCP mAb against HEK293 (1) and SMCP (AA: 1-116)-hIgGFc transfected HEK293 (2) cell lysate.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.