

SMC1 (phospho-Ser360) rabbit pAb

Cat No.: ES13059

For research use only

Overview

Product Name SMC1 (phospho-Ser360) rabbit pAb

Host species Rabbit

Applications WB;ELISA;IHC Species Cross-Reactivity Human;Mouse;Rat

Recommended dilutions WB 1:500-2000;IHC-p 1:50-300; ELISA 2000-20000 **Immunogen** Synthesized phosho peptide around human SMC1

(Ser360)

Specificity This antibody detects endogenous levels of

Human Mouse Rat SMC1 (phospho-Ser360)

Formulation Liquid in PBS containing 50% glycerol, 0.5% BSA and

0.02% sodium azide.

Storage Store at -20°C. Avoid repeated freeze-thaw cycles.

Protein Name SMC1 (Ser360)

Gene Name SMC1A DXS423E KIAA0178 SB1.8 SMC1 SMC1L1
Cellular localization Nucleus . Chromosome . Chromosome, centromere,

kinetochore. Associates with chromatin. Before prophase it is scattered along chromosome arms. During prophase, most of cohesin complexes dissociate from chromatin probably because of

phosphorylation by PLK,

Purification The antibody was affinity-purified from rabbit

antiserum by affinity-chromatography using

epitope-specific immunogen.

Clonality Polyclonal
Concentration 1 mg/ml
Observed band 143kD
Human Gene ID 8243
Human Swiss-Prot Number Q14683

Alternative Names Structural maintenance of chromosomes protein 1A

(SMC protein 1A) (SMC-1-alpha) (SMC-1A) (Sb1.8)

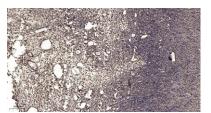
Background structural maintenance of chromosomes 1A(SMC1A)

Homo sapiens Proper cohesion of sister chromatids is a prerequisite for the correct





segregation of chromosomes during cell division. The cohesin multiprotein complex is required for sister chromatid cohesion. This complex is composed partly of two structural maintenance of chromosomes (SMC) proteins, SMC3 and either SMC1B or the protein encoded by this gene. Most of the cohesin complexes dissociate from the chromosomes before mitosis, although those complexes at the kinetochore remain. Therefore, the encoded protein is thought to be an important part of functional kinetochores. In addition, this protein interacts with BRCA1 and is phosphorylated by ATM, indicating a potential role for this protein in DNA repair. This gene, which belongs to the SMC gene family, is located in an area of the X-chromosome that escapes X inactivation. Mutations in this gene result in Cornelia de Lange syndrome. Altern



Immunohistochemical analysis of paraffin-embedded human oophoroma. 1, Antibody was diluted at 1:200(4° overnight). 2, Tris-EDTA,pH9.0 was used for antigen retrieval. 3,Secondary antibody was diluted at 1:200(room temperature, 45min).

