

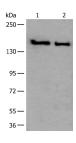
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SMC1A Polyclonal Antibody

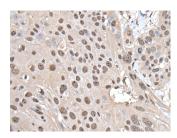
Catalog No.E-AB-52219ReactivityH,M,RStorageStore at -20°C. Avoid freeze / thaw cycles.HostRabbitApplicationsWB,IHC,ELISAIsotypeIgG

Note: Centrifuge before opening to ensure complete recovery of vial contents.

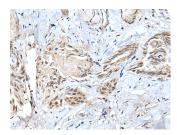
Images



Western blot analysis of HUVEC and Jurkat cell lysates using SMC1A Polyclonal Antibody at dilution of 1:300



Immunohistochemistry of paraffinembedded Human esophagus cancer tissue using SMC1A Polyclonal Antibody at dilution of 1:30(×200)



Immunohistochemistry of paraffinembedded Human thyroid cancer tissue using SMC1A Polyclonal Antibody at dilution of 1:30(×200)

Immunogen Information

Immunogen Fusion protein of human SMC1A

Gene Accession BC112127 **Swissprot** Q14683

Synonyms SMC protein 1A,SMC-1-alpha,SMC-1A,SMC1,SMC1

A,SMC1A,SMC1alpha,SMC1L1,SMCB

Product Information

Calculated MW 143 kDa

Observed MW Refer to figures

Buffer PBS with 0.05% NaN3 and 40% Glycerol,pH7.4

Purify Antigen affinity purification

Dilution WB 1:500-1:2000, IHC 1:25-1:100, ELISA

1:5000-1:10000

Background

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. Alternative splicing results in multiple transcript variants encoding different isoforms.

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