Resource Summary Report

Generated by FDI Lab - SciCrunch.org on Apr 4, 2025

SCP1 antibody

RRID:AB_301636 Type: Antibody

Proper Citation

(Abcam Cat# ab15090, RRID:AB_301636)

Antibody Information

URL: http://antibodyregistry.org/AB_301636

Proper Citation: (Abcam Cat# ab15090, RRID:AB_301636)

Target Antigen: SCP1 antibody

Host Organism: rabbit

Clonality: polyclonal

Comments: validation status unknown, seller recommendations provided in 2012: ICC/IF, IHC-Fr, IHC-P; Immunohistochemistry - frozen; Immunocytochemistry; Immunofluorescence; Immunohistochemistry; Immunohistochemistry - fixed

Antibody Name: SCP1 antibody

Description: This polyclonal targets SCP1 antibody

Target Organism: mouse, human

Antibody ID: AB_301636

Vendor: Abcam

Catalog Number: ab15090

Record Creation Time: 20241016T233820+0000

Record Last Update: 20241017T010109+0000

Ratings and Alerts

No rating or validation information has been found for SCP1 antibody.

No alerts have been found for SCP1 antibody.

Data and Source Information

Source: Antibody Registry

Usage and Citation Metrics

We found 21 mentions in open access literature.

Listed below are recent publications. The full list is available at FDI Lab - SciCrunch.org.

Biot M, et al. (2024) Principles of chromosome organization for meiotic recombination. Molecular cell, 84(10), 1826.

Chotiner JY, et al. (2024) TRIP13 localizes to synapsed chromosomes and functions as a dosage-sensitive regulator of meiosis. eLife, 12.

Shao Q, et al. (2023) ATF7IP2, a meiosis-specific partner of SETDB1, is required for proper chromosome remodeling and crossover formation during spermatogenesis. Cell reports, 42(8), 112953.

Zhang Q, et al. (2022) Homozygous Variant in KASH5 Causes Premature Ovarian Insufficiency by Disordered Meiotic Homologous Pairing. The Journal of clinical endocrinology and metabolism, 107(9), 2589.

Pereira C, et al. (2022) Multiple 9-1-1 complexes promote homolog synapsis, DSB repair, and ATR signaling during mammalian meiosis. eLife, 11.

Tanno N, et al. (2022) FBXO47 is essential for preventing the synaptonemal complex from premature disassembly in mouse male meiosis. iScience, 25(4), 104008.

Liu R, et al. (2021) YTHDC2 is essential for pachytene progression and prevents aberrant microtubule-driven telomere clustering in male meiosis. Cell reports, 37(11), 110110.

Du M, et al. (2021) PPP2R1B is modulated by ubiquitination and is essential for spermatogenesis. FASEB journal: official publication of the Federation of American Societies for Experimental Biology, 35(5), e21564.

Cheng EC, et al. (2021) The Essential Function of SETDB1 in Homologous Chromosome Pairing and Synapsis during Meiosis. Cell reports, 34(1), 108575.

Imai Y, et al. (2020) PRDM9 activity depends on HELLS and promotes local 5-

hydroxymethylcytosine enrichment. eLife, 9.

Abe H, et al. (2020) The Initiation of Meiotic Sex Chromosome Inactivation Sequesters DNA Damage Signaling from Autosomes in Mouse Spermatogenesis. Current biology: CB, 30(3), 408.

Shang Y, et al. (2020) MEIOK21: a new component of meiotic recombination bridges required for spermatogenesis. Nucleic acids research, 48(12), 6624.

Huang T, et al. (2020) The histone modification reader ZCWPW1 links histone methylation to PRDM9-induced double-strand break repair. eLife, 9.

Takemoto K, et al. (2020) Meiosis-Specific C19orf57/4930432K21Rik/BRME1 Modulates Localization of RAD51 and DMC1 to DSBs in Mouse Meiotic Recombination. Cell reports, 31(8), 107686.

Ishiguro KI, et al. (2020) MEIOSIN Directs the Switch from Mitosis to Meiosis in Mammalian Germ Cells. Developmental cell, 52(4), 429.

Gray S, et al. (2020) Cyclin N-Terminal Domain-Containing-1 Coordinates Meiotic Crossover Formation with Cell-Cycle Progression in a Cyclin-Independent Manner. Cell reports, 32(1), 107858.

DiTroia SP, et al. (2019) Maternal vitamin C regulates reprogramming of DNA methylation and germline development. Nature, 573(7773), 271.

Biswas U, et al. (2018) SMC1? Substitutes for Many Meiotic Functions of SMC1? but Cannot Protect Telomeres from Damage. Current biology: CB, 28(2), 249.

Diagouraga B, et al. (2018) PRDM9 Methyltransferase Activity Is Essential for Meiotic DNA Double-Strand Break Formation at Its Binding Sites. Molecular cell, 69(5), 853.

Zelazowski MJ, et al. (2017) Age-Dependent Alterations in Meiotic Recombination Cause Chromosome Segregation Errors in Spermatocytes. Cell, 171(3), 601.