

RAD50 Antibody

Purified Mouse Monoclonal Antibody Catalog # AO2232a

Product Information

Application WB, E **Primary Accession** Q92878 Reactivity Human Host Mouse Monoclonal Clonality **Clone Names** 5A8E5 Isotype IgG1 **Calculated MW** 153892

Description The protein encoded by this gene is highly similar to Saccharomyces

cerevisiae Rad50, a protein involved in DNA double-strand break repair. This protein forms a complex with MRE11 and NBS1. The protein complex binds to DNA and displays numerous enzymatic activities that are required for nonhomologous joining of DNA ends. This protein, cooperating with its partners, is important for DNA double-strand break repair, cell cycle checkpoint activation, telomere maintenance, and meiotic recombination. Knockout studies of the mouse homolog suggest this gene is essential for cell

growth and viability. Mutations in this gene are the cause of Nijmegen

breakage syndrome-like disorder.

Immunogen Purified recombinant fragment of human RAD50 (AA: 228-359) expressed in E.

Coli.

Formulation Purified antibody in PBS with 0.05% sodium azide

Additional Information

Gene ID 10111

Other Names DNA repair protein RAD50, hRAD50, 3.6.-.-, RAD50

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 RAD50 Antibody is for research use only and not for use in diagnostic or

therapeutic procedures.

Protein Information

Name

RAD50 {ECO:0000303 | PubMed:8756642, ECO:0000312 | HGNC:HGNC:9816}

Function

Component of the MRN complex, which plays a central role in double-strand break (DSB) repair, DNA recombination, maintenance of telomere integrity and meiosis (PubMed: 15064416, PubMed: 21757780, PubMed: 27889449, PubMed:28134932, PubMed:28867292, PubMed:9590181, PubMed:9651580, PubMed: 9705271). The MRN complex is involved in the repair of DNA double-strand breaks (DSBs) via homologous recombination (HR), an error-free mechanism which primarily occurs during S and G2 phases (PubMed: 15064416, PubMed: 21757780, PubMed: 27889449, PubMed: 28867292, PubMed: 9590181, PubMed: 9651580, PubMed: 9705271). The complex (1) mediates the end resection of damaged DNA, which generates proper single-stranded DNA, a key initial steps in HR, and is (2) required for the recruitment of other repair factors and efficient activation of ATM and ATR upon DNA damage (PubMed: 15064416, PubMed: 27889449, PubMed: <u>28867292</u>, PubMed: <u>9590181</u>, PubMed: <u>9651580</u>, PubMed: <u>9705271</u>). The MRN complex possesses single-strand endonuclease activity and double-strand-specific 3'-5' exonuclease activity, which are provided by MRE11, to initiate end resection, which is required for single-strand invasion and recombination (PubMed: 11741547, PubMed: 9590181, PubMed: 9651580, PubMed: 9705271). Within the complex, RAD50 is both required to bind DNA ends and hold them in close proximity and regulate the activity of MRE11 (PubMed:11741547, PubMed:12805565, PubMed:28134932). RAD50 provides an ATP-dependent control of MRE11 by positioning DNA ends into the MRE11 active site: ATP-binding induces a large structural change from an open form with accessible MRE11 nuclease sites into a closed form (By similarity). The MRN complex is also required for DNA damage signaling via activation of the ATM and ATR kinases: the nuclease activity of MRE11 is not required to activate ATM and ATR (PubMed: 15064416, PubMed: 15790808, PubMed: 16622404). The MRN complex is also required for the processing of R-loops (PubMed:31537797). In telomeres the MRN complex may modulate t-loop formation (PubMed: 10888888).

Cellular Location

Nucleus. Chromosome, telomere. Chromosome Note=Localizes to discrete nuclear foci after treatment with genotoxic agents (PubMed:10783165, PubMed:26215093). Localizes to DNA double- strand breaks (DSBs) (PubMed:15916964, PubMed:21757780)

Tissue Location

Expressed at very low level in most tissues, except in testis where it is expressed at higher level. Expressed in fibroblasts.

References

1.Breast Cancer Res Treat. 2010 Sep;123(2):607-9.2.Dis Markers. 2008;24(2):127-34.

Images

