

MRE11 Polyclonal Antibody

Catalog No: YT2829

Reactivity: Human; Mouse; Rat; Monkey

Applications: WB;IHC;IF;ELISA

Target: MRE11

Fields: >>Homologous recombination;>>Non-homologous end-joining;>>Cellular

senescence

P49959

Q61216

Gene Name: MRE11A

Protein Name: Double-strand break repair protein MRE11A

Human Gene Id: 4361

Human Swiss Prot

No:

Mouse Gene Id: 17535

Mouse Swiss Prot

No:

Rat Gene Id: 64046

Rat Swiss Prot No: Q9JIM0

Immunogen: The antiserum was produced against synthesized peptide derived from human

MRE11. AA range:230-279

Specificity: MRE11 Polyclonal Antibody detects endogenous levels of MRE11 protein.

Formulation : Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Source: Polyclonal, Rabbit, IgG

Dilution : WB 1:500 - 1:2000. IHC 1:100 - 1:300. ELISA: 1:20000.. IF 1:50-200

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Purification: The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Concentration: 1 mg/ml

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Observed Band: 80kD

Cell Pathway: Homologous recombination; Non-homologous end-joining;

Background: This gene encodes a nuclear protein involved in homologous recombination,

telomere length maintenance, and DNA double-strand break repair. By itself, the protein has 3' to 5' exonuclease activity and endonuclease activity. The protein forms a complex with the RAD50 homolog; this complex is required for nonhomologous joining of DNA ends and possesses increased single-stranded DNA endonuclease and 3' to 5' exonuclease activities. In

conjunction with a DNA ligase, this protein promotes the joining of

noncomplementary ends in vitro using short homologies near the ends of the DNA fragments. This gene has a pseudogene on chromosome 3. Alternative splicing of this gene results in two transcript variants encoding different isoforms. [provided

by RefSeq, Jul 2008],

Function: cofactor:Manganese.,disease:Defects in MRE11A are a cause of ataxia

telangiectasia-like disorder (ATLD) [MIM:604391]. ATLD is a disease with the same clinical feature than ataxia-telangiectasia but with a somewhat milder clinical course., disease:Defects in MRE11A may be a cause of breast cancer., 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. The complex possesses single-strand endonuclease activity and double-strand-specific 3'-5' exonuclease activity, which are provided by MRE11A. RAD50 may be required to bind DNA ends and hold them in close proximity. This could facilitate searches for short or long regions of sequence

of DNA ligases and/or restrict the nuclease activity of MRE11A to prev

Subcellular Location:

Nucleus . Chromosome, telomere . Chromosome . Localizes to discrete nuclear

homology in the recombining DNA templates, and may also stimulate the activity

foci after treatment with genotoxic agents. .

Expression: Bladder, Brain, Epithelium, Lung,

Tag: orthogonal

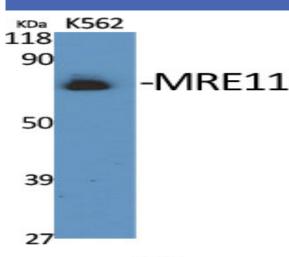
Sort : 10207



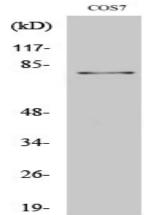
Host: Rabbit

Modifications: Unmodified

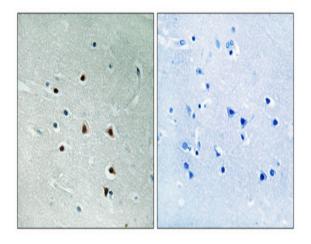
Products Images



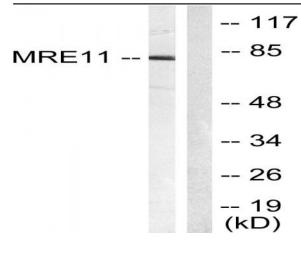
Western Blot analysis of various cells using MRE11 Polyclonal Antibody cells nucleus extracted by Minute TM Cytoplasmic and Nuclear Fractionation kit (SC-003,Inventbiotech,MN,USA).



Western Blot analysis of COS7 cells using MRE11 Polyclonal Antibody cells nucleus extracted by Minute TM Cytoplasmic and Nuclear Fractionation kit (SC-003,Inventbiotech,MN,USA).



Immunohistochemical analysis of paraffin-embedded Human brain. Antibody was diluted at 1:100(4° overnight). High-pressure and temperature Tris-EDTA,pH8.0 was used for antigen retrieval. Negetive contrl (right) obtaned from antibody was pre-absorbed by immunogen peptide.



Western blot analysis of lysates from Jurkat cells, treated with UV 15', using MRE11 Antibody. The lane on the right is blocked with the synthesized peptide.