

XRCC4 Monoclonal Antibody(5C10)

Catalog # AP63340

Product Information

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|-------------------|------------------------|
| Application | WB, IHC-P, IF, IP |
| Primary Accession | Q13426 |
| Reactivity | Human |
| Host | Mouse |
| Clonality | Monoclonal |
| Calculated MW | 38287 |

Additional Information

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| Gene ID | 7518 |
| Other Names | XRCC4; DNA repair protein XRCC4; X-ray repair cross-complementing protein 4 |
| Dilution | WB~~WB: 1:2000 IP:1:200 IF 1:200 IHC 1:50-300 IHC-P~~N/A IF~~1:50~200 IP~~N/A |
| Format | PBS, pH 7.4, containing 0.09% (W/V) sodium azide as Preservative and 50% Glycerol. |
| Storage Conditions | -20°C |

Protein Information

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| Name | XRCC4 {ECO:0000303 PubMed:8548796, ECO:0000312 HGNC:HGNC:12831} |
| Function | [DNA repair protein XRCC4]: DNA non-homologous end joining (NHEJ) core factor, required for double-strand break repair and V(D)J recombination (PubMed: 10757784 , PubMed: 10854421 , PubMed: 12517771 , PubMed: 16412978 , PubMed: 17124166 , PubMed: 17290226 , PubMed: 22228831 , PubMed: 25597996 , PubMed: 25742519 , PubMed: 25934149 , PubMed: 26100018 , PubMed: 26774286 , PubMed: 8548796). Acts as a scaffold protein that regulates recruitment of other proteins to DNA double-strand breaks (DSBs) (PubMed: 15385968 , PubMed: 20852255 , PubMed: 26774286 , PubMed: 27437582). Associates with NHEJ1/XLF to form alternating helical filaments that bridge DNA and act like a bandage, holding together the broken DNA until it is repaired (PubMed: 21768349 , PubMed: 21775435 , PubMed: 22287571 , PubMed: 26100018 , PubMed: 27437582 , PubMed: 28500754). The XRCC4-NHEJ1/XLF subcomplex binds to the DNA fragments of a DSB in a highly diffusive manner and robustly bridges two independent DNA molecules, holding the broken DNA fragments in close proximity to one other (PubMed: 27437582). The mobility of the bridges ensures that the ends remain |

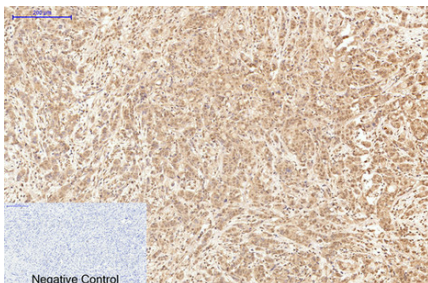
accessible for further processing by other repair factors (PubMed:[27437582](#)). Plays a key role in the NHEJ ligation step of the broken DNA during DSB repair via direct interaction with DNA ligase IV (LIG4): the LIG4-XRCC4 subcomplex reseals the DNA breaks after the gap filling is completed (PubMed:[10757784](#), PubMed:[10854421](#), PubMed:[12517771](#), PubMed:[17290226](#), PubMed:[19837014](#), PubMed:[9242410](#)). XRCC4 stabilizes LIG4, regulates its subcellular localization and enhances LIG4's joining activity (PubMed:[10757784](#), PubMed:[10854421](#), PubMed:[12517771](#), PubMed:[17290226](#), PubMed:[21982441](#), PubMed:[22228831](#), PubMed:[9242410](#)). Binding of the LIG4-XRCC4 subcomplex to DNA ends is dependent on the assembly of the DNA-dependent protein kinase complex DNA-PK to these DNA ends (PubMed:[10757784](#), PubMed:[10854421](#)). Promotes displacement of PNKP from processed strand break termini (PubMed:[20852255](#), PubMed:[28453785](#)).

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|--------------------------|--|
| Cellular Location | Nucleus. Chromosome. Note=Localizes to site of double-strand breaks. |
| Tissue Location | Widely expressed.. |

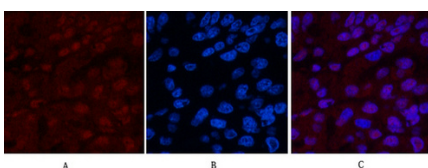
Background

Involved in DNA non-homologous end joining (NHEJ) required for double-strand break repair and V(D)J recombination. Binds to DNA and to DNA ligase IV (LIG4). The LIG4-XRCC4 complex is responsible for the NHEJ ligation step, and XRCC4 enhances the joining activity of LIG4. Binding of the LIG4-XRCC4 complex to DNA ends is dependent on the assembly of the DNA-dependent protein kinase complex DNA-PK to these DNA ends.

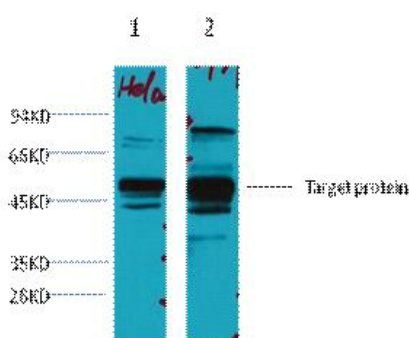
Images



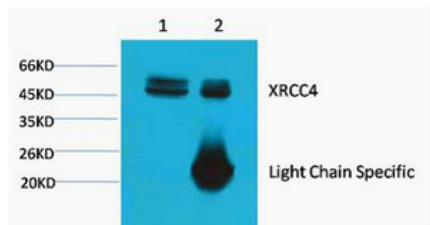
Immunohistochemical analysis of paraffin-embedded Human-breast-cancer tissue. 1,XRCC4 Monoclonal Antibody(5C10) was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room temperature, 30min). Negative control was used by secondary antibody only.



Immunofluorescence analysis of Human-liver-cancer tissue. 1,XRCC4 Monoclonal Antibody(5C10)(red) was diluted at 1:200(4°C,overnight). 2, Cy3 labeled Secondary antibody was diluted at 1:300(room temperature, 50min).3, Picture B: DAPI(blue) 10min. Picture A:Target. Picture B: DAPI. Picture C: merge of A+B



Western blot analysis of 1) HeLa, 2) 293T, diluted at 1:3000.



1) Input: Hela Cell Lysate 2) IP product: IP dilute 1:200

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.