Product datasheet

Anti-BRCA2 antibody ab123491

Overview

Product name Anti-BRCA2 antibody
Description Rabbit polyclonal to BRCA2
Host species Rabbit
Tested applications Suitable for: WB, IP
Species reactivity Reacts with: Human
Predicted to work with: Chimpanzee, Rhesus monkey, Gorilla, Orangutan
Immunogen Synthetic peptide, corresponding to a region within amino acids 450-500 of Human BRCA2 using the numbering given in entry NP_000050.1.
Positive control Whole cell lysate from 293T, HeLa and Jurkat cells

Properties

Form Liquid
Storage instructions Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.
Storage buffer Preservative: 0.09% Sodium azide
Constituent: 99% Tris citrate/phosphate
pH 7 to 8
Purity Immunogen affinity purified
Clonality Polyclonal
Isotype IgG

Applications

Our Abpromise guarantee covers the use of ab123491 in the following tested applications.
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

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### Function
Involved in double-strand break repair and/or homologous recombination. Binds RAD51 and potentiates recombinational DNA repair by promoting assembly of RAD51 onto single-stranded DNA (ssDNA). Acts by targeting RAD51 to ssDNA over double-stranded DNA, enabling RAD51 to displace replication protein-A (RPA) from ssDNA and stabilizing RAD51-ssDNA filaments by blocking ATP hydrolysis. May participate in S phase checkpoint activation. Binds selectively to ssDNA, and to ssDNA in tailed duplexes and replication fork structures.

### Tissue specificity
Highest levels of expression in breast and thymus, with slightly lower levels in lung, ovary and spleen.

### Involvement in disease
Defects in BRCA2 are a cause of susceptibility to breast cancer (BC) [MIM:114480]. A common malignancy originating from breast epithelial tissue. Breast neoplasms can be distinguished by their histologic pattern. Invasive ductal carcinoma is by far the most common type. Breast cancer is etiologically and genetically heterogeneous. Important genetic factors have been indicated by familial occurrence and bilateral involvement. Mutations at more than one locus can be involved in different families or even in the same case.

Defects in BRCA2 are the cause of pancreatic cancer type 2 (PNCA2) [MIM:613347]. It is a malignant neoplasm of the pancreas. Tumors can arise from both the exocrine and endocrine portions of the pancreas, but 95% of them develop from the exocrine portion, including the ductal epithelium, acinar cells, connective tissue, and lymphatic tissue.

Defects in BRCA2 are a cause of susceptibility to breast-ovarian cancer familial type 2 (BROVCA2) [MIM:612555]. A condition associated with familial predisposition to cancer of the breast and ovaries. Characteristic features in affected families are an early age of onset of breast cancer (often before age 50), increased chance of bilateral cancers (cancer that develop in both breasts, or both ovaries, independently), frequent occurrence of breast cancer among men, increased incidence of tumors of other specific organs, such as the prostate.

Defects in BRCA2 are the cause of Fanconi anemia complementation group D type 1 (FANCD1) [MIM:605724]. It is a disorder affecting all bone marrow elements and resulting in anemia, leukopenia and thrombopenia. It is associated with cardiac, renal and limb malformations, dermal pigmentary changes, and a predisposition to the development of malignancies. At the cellular level it is associated with hypersensitivity to DNA-damaging agents, chromosomal instability (increased chromosome breakage) and defective DNA repair.

Defects in BRCA2 are a cause of glioma type 3 (GLM3) [MIM:613029]. Gliomas are benign or malignant central nervous system neoplasms derived from glial cells. They comprise astrocytomas and glioblastoma multiforme that are derived from astrocytes, oligodendrogliomas derived from oligodendrocytes and ependymomas derived from ependymocytes.

### Sequence similarities
Contains 8 BRCA2 repeats.

### Post-translational modifications
Phosphorylated by ATM upon irradiation-induced DNA damage. Ubiquitinated in the absence of DNA damage; this does not lead to proteasomal degradation. In contrast, ubiquitination in response to DNA damage leads to proteasomal degradation.

### Images

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<td>IP</td>
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<td>Use at 2-10 µg/mg of lysate.</td>
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All lanes: Anti-BRCA2 antibody (ab123491) at 0.1 µg/ml

Lane 1: 293T whole cell lysate at 50 µg
Lane 2: 293T whole cell lysate at 15 µg
Lane 3: 293T whole cell lysate at 5 µg
Lane 5: HeLa whole cell lysate at 50 µg
Lane 6: HeLa whole cell lysate at 15 µg

Predicted band size: 384 kDa

Exposure time: 30 seconds

Detection of Human BRCA2 by Immunoprecipitation from HeLa whole cell lysate (1 mg for IP, 20% of IP loaded), using ab123491 at 6 µg/mg lysate.
Subsequent Western blot detection used ab123491 at 1 µg/ml.
Detection: Chemiluminescence with an exposure time of 10 seconds.

All lanes: Anti-BRCA2 antibody (ab123491) at 0.1 µg/ml

Lane 1: Whole cell lysate from HeLa at 50 µg/ml
Lane 2: Whole cell lysate from HeLa at 15 µg/ml
Lane 3: Whole cell lysate from 293T at 50 µg/ml
Lane 4: Whole cell lysate from Jurkat at 50 µg/ml

Developed using the ECL technique.

Predicted band size: 384 kDa

Exposure time: 10 minutes

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