

## **Anti-BRCA1** antibody



**Description** Rabbit polyclonal to BRCA1.

Model STJ91882

**Host** Rabbit

**Reactivity** Human

**Applications** ELISA, IHC

**Immunogen** Synthesized peptide derived from human BRCA1 around the non-

phosphorylation site of S988.

**Immunogen Region** 930-1010 aa

**Gene ID** <u>672</u>

Gene Symbol BRCA1

**Dilution range** IHC 1:100-1:300ELISA 1:20000

**Specificity** BRCA1 Polyclonal Antibody detects endogenous levels of BRCA1 protein.

**Tissue Specificity** Isoform 1 and isoform 3 are widely expressed. Isoform 3 is reduced or absent

in several breast and ovarian cancer cell lines.

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

chromatography using epitope-specific immunogen.

**Note** For Research Use Only (RUO).

**Protein Name**Breast cancer type 1 susceptibility protein RING finger protein 53 RING-type

E3 ubiquitin transferase BRCA1

Molecular Weight 180 kDa

**Clonality** Polyclonal

**Conjugation** Unconjugated

**Isotype** IgG

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

**Concentration** 1 mg/ml

**Storage Instruction** Store at -20°C, and avoid repeat freeze-thaw cycles.

Database Links <u>HGNC:1100OMIM:113705</u>

**Alternative Names** Breast cancer type 1 susceptibility protein RING finger protein 53 RING-type

E3 ubiquitin transferase BRCA1

**Function** E3 ubiquitin-protein ligase that specifically mediates the formation of 'Lys-6'-

linked polyubiquitin chains and plays a central role in DNA repair by facilitating cellular responses to DNA damage. It is unclear whether it also mediates the formation of other types of polyubiquitin chains. The E3 ubiquitin-protein ligase activity is required for its tumor suppressor function. The BRCA1-BARD1 heterodimer coordinates a diverse range of cellular pathways such as DNA damage repair, ubiquitination and transcriptional regulation to maintain genomic stability. Regulates centrosomal microtubule nucleation. Required for normal cell cycle progression from G2 to mitosis. Required for appropriate cell cycle arrests after ionizing irradiation in both the S-phase and the G2 phase of the cell cycle. Involved in transcriptional

regulation of P21 in response to DNA damage. Required for FANCD2 targeting to sites of DNA damage. May function as a transcriptional regulator. Inhibits lipid synthesis by binding to inactive phosphorylated ACACA and

repair (HRR) via its direct interaction with PALB2, fine-tunes

recombinational repair partly through its modulatory role in the PALB2-dependent loading of BRCA2-RAD51 repair machinery at DNA breaks. Component of the BRCA1-RBBP8 complex which regulates CHEK1 activation and controls cell cycle G2/M checkpoints on DNA damage via BRCA1-mediated ubiquitination of RBBP8. Acts as a transcriptional activator

preventing its dephosphorylation. Contributes to homologous recombination

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**Sequence and Domain Family** The BRCT domains recognize and bind phosphorylated pSXXF motif on

proteins. The interaction with the phosphorylated pSXXF motif of ABRAXAS1, recruits BRCA1 at DNA damage sites. The RING-type zinc

finger domain interacts with BAP1.

Cellular Localization Nucleus Chromosome Cytoplasm. Localizes at sites of DNA damage at

double-strand breaks (DSBs). recruitment to DNA damage sites is mediated by ABRAXAS1 and the BRCA1-A complex. Translocated to the cytoplasm during UV-induced apoptosis. Isoform 3: Cytoplasm. Isoform 5: Cytoplasm

Post-translational

**Modifications** 

Phosphorylation at Ser-308 by AURKA is required for normal cell cycle progression from G2 to mitosis. Phosphorylated in response to IR, UV, and various stimuli that cause checkpoint activation, probably by ATM or ATR. Phosphorylation at Ser-988 by CHEK2 regulates mitotic spindle assembly. Autoubiquitinated, undergoes 'Lys-6'-linked polyubiquitination. 'Lys-6'-linked

polyubiquitination does not promote degradation.

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