Phospho-BRCA1 (Ser1524) Ab

Cat.#: AF3289 Concn.: 1mg/ml Mol.Wt.: 207 kDa Size: 100ul.200ul Source: Rabbit Clonality: Polyclonal

Application: WB 1:500-1:2000 IHC 1:50-1:200

Reactivity: Human, Mouse, Rat

Purification: The Ab is from purified rabbit serum by affinity purification

via seguential chromatography on phospho- and non-

phospho-peptide affinity columns.

Specificity: Phospho-BRCA1 (Ser1524) Ab detects endogenous levels of

BRCA1 only when phosphorylated at Serine 1524.

A synthesized peptide derived from human BRCA1 around Immunogen:

the phosphorylation site of Serine 1524.

Uniprot: P38398

Description: This gene encodes a nuclear phosphoprotein that plays a

> role in maintaining genomic stability and acts as a tumor suppressor. The encoded protein combines with other tumor suppressors, DNA damage sensors, and signal transducers to form a large multi-subunit protein complex known as BASC for BRCA1-associated genome surveillance complex.

Subcellular Location: Cytoplasm; Nucleus. Localizes at sites of DNA damage at

> double-strand breaks (DSBs) and recruitment to DNA damage sites is mediated by the BRCA1-A complex.

Tissue Specificity: Isoform 1 and isoform 3 are widely expressed. Isoform 3 is

reduced or absent in several breast and ovarian cancer cell

lines.

Similarity: 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.

Storage Condition and

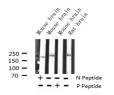
Buffer:

Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Store at -20

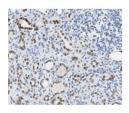
°C.Stable for 12 months from date of receipt.



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Western blot analysis of Phospho-BRCA1 (Ser1524) Ab expression in mouse brain and rat brain tissues lysates.



AF3289 at 1/100 staining human Kidney carcinoma tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue was then blocked and incubated with the Ab for 1.5 hours at 22°C. An HRP conjugated goat antirabbit Ab was used as the secondary.

<code>IMPORTANT:</code> For western blot, incubate membrane with diluted primary Ab in 5% w/v milk , 1X TBS, 0.1% Tween®20 at 4°C with gentle shaking, overnight.

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