



FANCD2 (phospho Ser222) rabbit pAb

Cat#: orb768070 (Manual)

For research use only. Not intended for diagnostic use.

Product Name FANCD2 (phospho Ser222) rabbit pAb

Host species Rabbit

Applications WB;IHC;IF;ELISA

Species Cross-Reactivity Human; Mouse; Rat

Recommended dilutions Western Blot: 1/500 - 1/2000. Immunohistochemistry: 1/100 - 1/300. ELISA:

1/5000. Not yet tested in other applications.

Immunogen The antiserum was produced against synthesized peptide derived from

human FANCD2 around the phosphorylation site of Ser222. AA range:188-

237

Specificity Phospho-FANCD2 (S222) Polyclonal Antibody detects endogenous levels of

FANCD2 protein only when phosphorylated at S222.

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

azide..

Storage Store at -20°C. Avoid repeated freeze-thaw cycles.

Protein Name Fanconi anemia group D2 protein

Gene Name FANCD2

Cellular localization Nucleus. Concentrates in nuclear foci during S phase and upon genotoxic

stress. At the onset of mitosis, excluded from chromosomes and diffuses into the cytoplasm, returning to the nucleus at the end of cell division. Observed

in a few spots localized in

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

chromatography using epitope-specific immunogen.





Clonality Polyclonal

Concentration 1 mg/ml

Observed band 166kD

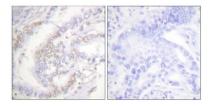
Human Gene ID 2177

Human Swiss-Prot Number Q9BXW9

Alternative Names FANCD2; FACD; Fanconi anemia group D2 protein; Protein FACD2

Background

Fanconi anemia complementation group D2(FANCD2) Homo sapiens The Fanconi anemia complementation group (FANC) currently includes FANCA, FANCB, FANCC, FANCD1 (also called BRCA2), FANCD2, FANCE, FANCF, FANCG, FANCI, FANCJ (also called BRIP1), FANCL, FANCM and FANCN (also called PALB2). The previously defined group FANCH is the same as FANCA. Fanconi anemia is a genetically heterogeneous recessive disorder characterized by cytogenetic instability, hypersensitivity to DNA crosslinking agents, increased chromosomal breakage, and defective DNA repair. The members of the Fanconi anemia complementation group do not share sequence similarity; they are related by their assembly into a common nuclear protein complex. This gene encodes the protein for complementation group D2. This protein is monoubiquinated in response to DNA damage, resulting in its localization to nuclear foci with other proteins (BRCA1 AND BRCA2) involved in homology-directed DNA renai

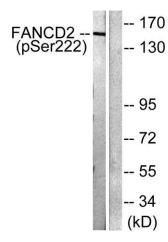


Immunohistochemistry analysis of paraffin-embedded human lung carcinoma, using FANCD2 (Phospho-Ser222) Antibody. The picture on the right is blocked with the phospho peptide.





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Western blot analysis of lysates from HT29 cells treated with Calyculin A 50ng/ml 30', using FANCD2 (Phospho-Ser222) Antibody. The lane on the right is blocked with the phospho peptide.