

FANCA (Phospho Ser1149) Rabbit pAb

CatalogNo: YP0943 Orthogonal Validated 

Key Features

Host Species

- Rabbit

Reactivity

- Human, Mouse, Rat

Applications

- WB, IHC, IF, ELISA

MW

- 162kD (Observed)

Isotype

- IgG

Storage

Storage* -15°C to -25°C/1 year (Do not lower than -25°C)

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

Recommended Dilution Ratios

WB 1:500-1:2000

IHC 1:100-1:300

IF 1:200-1:1000

ELISA 1:5000

Not yet tested in other applications.

Basic Information

Clonality Polyclonal

Immunogen Information

Immunogen The antiserum was produced against synthesized peptide derived from human FANCA around the phosphorylation site of Ser1149. AA range: 1121-1170

Specificity

Phospho-FANCA (S1149) Polyclonal Antibody detects endogenous levels of FANCA protein only when phosphorylated at S1149. The name of modified sites may be influenced by many factors, such as species (the modified site was not originally found in human samples) and the change of protein sequence (the previous protein sequence is incomplete, and the protein sequence may be prolonged with the development of protein sequencing technology). When naming, we will use the "numbers" in historical reference to keep the sites consistent with the reports. The antibody binds to the following modification sequence (lowercase letters are modification sites):DPsLM

| Target Information

Gene name FANCA

Protein Name Fanconi anemia group A protein

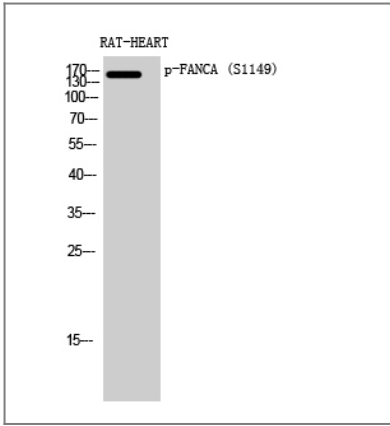
Organism	Gene ID	UniProt ID
Human	2175;	O15360;
Mouse		Q9JL70;

Cellular Localization Nucleus. Cytoplasm. The major form is nuclear. The minor form is cytoplasmic.

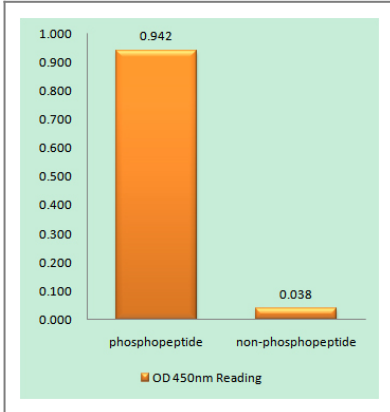
Tissue specificity Cervix, Epithelium, Lymphoblast, Ovary, PCR rescued clones,

Function Disease: Defects in FANCA are a cause of Fanconi anemia (FA) [MIM:227650]. FA is a genetically heterogeneous, autosomal recessive disorder characterized by progressive pancytopenia, a diverse assortment of congenital malformations, 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. Function: DNA repair protein that may operate in a postreplication repair or a cell cycle checkpoint function. May be involved in interstrand DNA cross-link repair and in the maintenance of normal chromosome stability. PTM: Phosphorylated upon DNA damage, probably by ATM or ATR. Phosphorylation is required for the formation of the nuclear complex. Not phosphorylated in cells derived from groups A, B, C, E, F, G, and H. subcellular location: The major form is nuclear. The minor form is cytoplasmic. subunit: Belongs to the multisubunit FA complex composed of FANCA, FANCB, FANCC, FANCE, FANCF, FANCG, FANCL/PHF9 and FANCM. The complex is not found in FA patients.

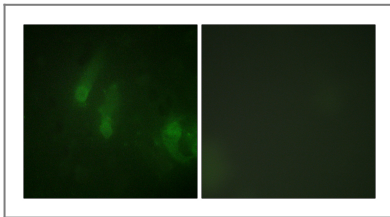
| Validation Data



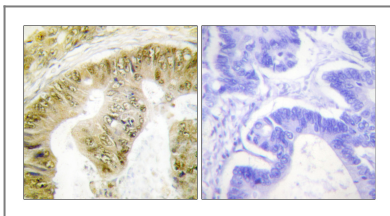
Western Blot analysis of RAT-HEART cells using Phospho-FANCA (S1149) Polyclonal Antibody diluted at 1:500



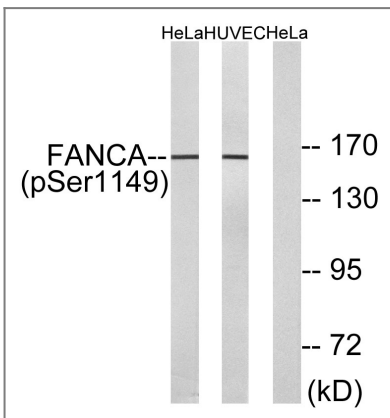
Enzyme-Linked Immunosorbent Assay (Phospho-ELISA) for Immunogen Phosphopeptide (Phospho-left) and Non-Phosphopeptide (Phospho-right), using FANCA (Phospho-Ser1149) Antibody



Immunofluorescence analysis of HeLa cells, using FANCA (Phospho-Ser1149) Antibody. The picture on the right is blocked with the phospho peptide.



Immunohistochemistry analysis of paraffin-embedded human colon carcinoma, using FANCA (Phospho-Ser1149) Antibody. The picture on the right is blocked with the phospho peptide.



Western blot analysis of lysates from HeLa cells treated with IGF 100ng/ml 10' and HUVEC cells treated with EGF 200ng/ml 30', using FANCA (Phospho-Ser1149) Antibody. The lane on the right is blocked with the phospho peptide.

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Please scan the QR code to access additional product information:
FANCA (Phospho Ser1149) Rabbit pAb

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