## FANCD2 Rabbit Polyclonal Antibody

Catalog No: #54616

Package Size: #54616-1 50ul #54616-2 100ul



Orders: order@signalwayantibody.com Support: tech@signalwayantibody.com

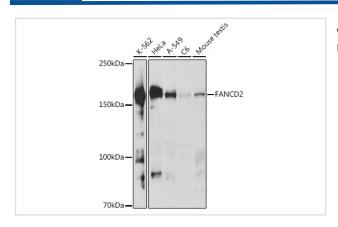
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Product Name	FANCD2 Rabbit Polyclonal Antibody	
Host Species	Rabbit	
Clonality	Polyclonal	
Isotype	IgG	
Purification	Affinity purification	
Applications	WB,IF	
Species Reactivity	Human,Mouse,Rat	
Immunogen Description	Recombinant protein of human FANCD2.	
Other Names	FA4;FAD;FACD;FAD2;FANCD;FANCD2	
Accession No.	Swiss Prot:Q9BXW9GeneID:2177	
Uniprot	Q9BXW9	
Calculated MW	164kDa	
SDS-PAGE MW	164kDa	
Formulation	Buffer: PBS with 0.02% sodium azide,50% glycerol,pH7.3.	
Storage	Store at -20°C. Avoid freeze / thaw cycles.	

## **Application Details**

WB□1:500 - 1:2000IF□1:50 - 1:200

## **Images**



Western blot analysis of extracts of various cell lines, using FANCD2 at 1:1000 dilution.

## Background

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 repair. Alternative splicing results in multiple transcript variants.

Note: This product is for in vitro research use only