Elabscience®

FANCD2 Polyclonal Antibody

catalog number: E-AB-60589

Note: Centrifuge before opening to ensure complete recovery of vial contents.

Description			
Reactivity	Human		
Immunogen	Recombinant fusion protein of human FANCD2 (NP_149075.2).		
Host	Rabbit	Rabbit	
Isotype	IgG		
Purification	Affinity purification	Affinity purification	
Buffer	Phosphate buffered solu	Phosphate buffered solution, pH 7.4, containing 0.05% stabilizer and 50% glycerol.	
Applications	Recommended Dilution		
IF	1:50-1:200	1:50-1:200	
Data			
Immunofluorescence analysis of MCF-7 cells using FANCD2 Immunofluorescence analysis of GFP-RNF168 transg		2 Immunofluorescence analysis of GFP-RNF168 transgenic	
Poly	clonal Antibody	U2OS cells using FANCD2 Polyclonal Antibody	
Preparation & Storage			
Storage	Store at -20°C Valid for 1	Store at -20°C Valid for 12 months. Avoid freeze / thaw cycles.	
Shipping	The product is shipped v	The product is shipped with ice pack, upon receipt, store it immediately at the	
	temperature recommended.		
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.

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