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FANCM Rabbit pAb

Catalog No.: A7602

Basic Information

Observed MW

Refer to figures

Calculated MW

75kDa/229kDa/232kDa

Category

Primary antibody

Applications

WB,IF/ICC

Cross-Reactivity

Human

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 M. Alternative splicing results in multiple transcript variants.

Recommended Dilutions

WB 1:500 - 1:2000

IF/ICC 1:50 - 1:100

Immunogen Information

Gene ID57697

Swiss Prot
Q8IYD8

Immunogen

Recombinant fusion protein containing a sequence corresponding to amino acids 390-660 of human FANCM (NP_065988.1).

Synonyms

FANCM; FAAP250; KIAA1596

Contact

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Product Information

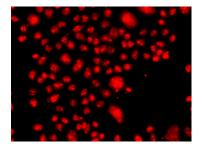
SourceIsotypePurificationRabbitIgGAffinity purification

Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.02% sodium azide,50% glycerol,pH7.3.

Validation Data



Immunofluorescence analysis of A549 cells using FANCM antibody (A7602).