



# FANCB mouse mAb

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|--------------------|--|
| Catalog No         | BYmab-12189  |
| Isotype            | IgG  |
| Reactivity         | Human; Mouse   |
| Applications       | WB   |
| Gene Name          | FANCB  |
| Protein Name       | FANCB  |
| Immunogen          | Synthesized peptide derived from human FANCB AA range: 585-635   |
| Specificity        | This antibody detects endogenous levels of FANCB at Human/Mouse  |
| Formulation        | Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.  |
| Source             | Monoclonal, Mouse,IgG  |
| Purification       | The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.   |
| Dilution           | WB 1:500-2000  |
| Concentration      | 1 mg/ml  |
| Purity             | ≥90%   |
| Storage Stability  | -20°C/1 year   |
| Synonyms           |  |
| Observed Band      |  |
| Cell Pathway       | Nucleus .  |
| Tissue Specificity |  |
| Function           | disease:Defects in FANCB 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.,disease:Defects in FANCB are the cause of Fanconi anemia complementation group B (FANCB) [MIM:300514]; also called Fanconi pancytopenia type 2 (FA2).,disease:Defects in FANCB are the cause of X-linked VACTERL-H (XVACTERL-H) [MIM:314390]; also known as X-linked VACTERL association with hydrocephalus syndrome. VACTERL is an acronym for vertebral anomalies, anal atresia, cardiac malformations, tracheoesophageal fistula, renal |

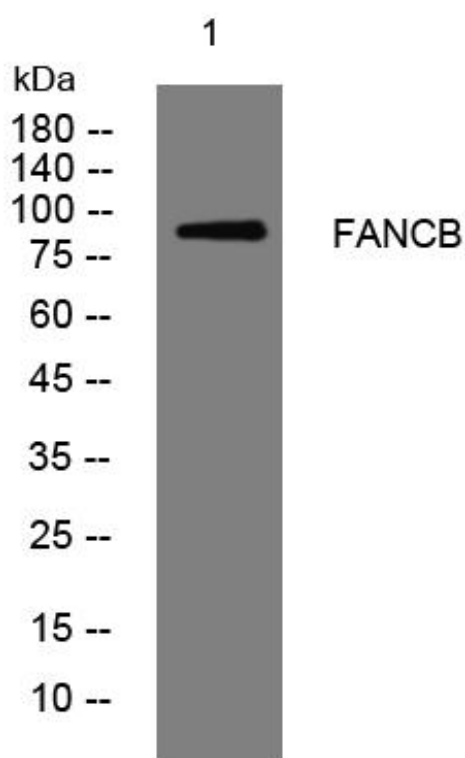
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anomalie

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|----------------------------------|--|
| <b>Background</b>                | This gene encodes a member of the Fanconi anemia complementation group B. This protein is assembled into a nucleoprotein complex that is involved in the repair of DNA lesions. Mutations in this gene can cause chromosome instability and VACTERL syndrome with hydrocephalus. [provided by RefSeq, Apr 2016], |
| <b>matters needing attention</b> | Avoid repeated freezing and thawing!   |
| <b>Usage suggestions</b>         | This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.  |

## Products Images



Western Blot analysis of various cells using FANCB mouse mAb

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