

Anti-Zebrafish FANCM Antibody Picoband®

Catalog Number: AZA8QLH8

About FANCM

Predicted to enable 3'-5' DNA helicase activity; four-way junction DNA binding activity; and four-way junction helicase activity. Predicted to be involved in double-strand break repair via synthesis-dependent strand annealing and interstrand cross-link repair. Predicted to act upstream of or within DNA repair. Predicted to be located in nucleus. Is expressed in central nervous system; eye; and lens. Human ortholog(s) of this gene implicated in primary ovarian insufficiency 15 and spermatogenic failure 28. Orthologous to human FANCM (FA complementation group M).

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Overview

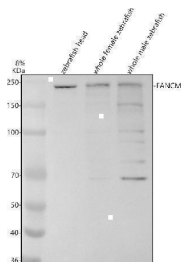
Product Name	Anti-Zebrafish FANCM Antibody Picoband®
Reactive Species	Zebrafish
Description	Boster Bio Anti-Zebrafish FANCM Antibody Picoband® catalog # AZA8QLH8. Tested in WB applications. This antibody reacts with Zebrafish. The brand Picoband indicates this is a premium antibody that guarantees superior quality, high affinity, and strong signals with minimal background in Western blot applications. Only our best-performing antibodies are designated as Picoband, ensuring unmatched performance.
Application	WB
Clonality	Polyclonal
Formulation	Each vial contains 4 mg Trehalose, 0.9 mg NaCl, 0.2 mg Na ₂ HPO ₄ .
Storage Instructions	At -20°C for one year from date of receipt. After reconstitution, at 4°C for one month. It can also be aliquotted and stored frozen at -20°C for six months. Avoid repeated freezing and thawing.
Host	Rabbit
Uniprot ID	A8QLH8

Technical Details

Immunogen	E.coli-derived Zebrafish FANCM recombinant protein (Position: D222-Q1491).
Form	Lyophilized

Concentration	Adding 0.2 ml of distilled water will yield a concentration of 500 ug/ml.
Purification	Immunogen affinity purified.
Suggested Dilutions	Western blot, 0.25-0.5 ug/ml, Zebrafish

Anti-Zebrafish FANCM Antibody Picoband® (AZA8QLH8) Images



Western blot analysis of FANCM using anti-FANCM antibody (AZA8QLH8). Electrophoresis was performed on a 8% SDS-PAGE gel at 80V (Stacking gel) / 120V (Resolving gel) for 2 hours. The sample well of each lane was loaded with 30 ug of sample under reducing conditions. Lane 1: zebrafish head tissue lysates, Lane 2: whole female zebrafish tissue lysates, Lane 3: whole male zebrafish tissue lysates. After electrophoresis, proteins were transferred to a nitrocellulose membrane at 150 mA for 50-90 minutes. Blocked the membrane with 5% non-fat milk/TBS for 1.5 hour at RT. The membrane was incubated with rabbit anti-FANCM antigen affinity purified polyclonal antibody (AZA8QLH8) at 0.5 ug/mL overnight at 4°C, then washed with TBS-0.1%Tween 3 times with 5 minutes each and probed with a goat anti-rabbit IgG-HRP secondary antibody at a dilution of 1:5000 for 1.5 hour at RT. The signal is developed using an ECL Plus Western Blotting Substrate (Catalog # AR1196-200) with Tanon 5200 system. A specific band was detected for FANCM at approximately 232 kDa. The expected band size for FANCM is at 229 kDa.

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For Research Use Only. Not for use in diagnostic procedures.