

Anti-Zebrafish FOXA1 Antibody Picoband®

Catalog Number: AZQ6POC9

About FOXA1

Enables transcription cis-regulatory region binding activity. Acts upstream of or within axial mesoderm morphogenesis. Predicted to be located in nucleus. Is expressed in several structures, including central nervous system; chordo neural hinge; digestive system; endoderm; and liver primordium. Human ortholog(s) of this gene implicated in breast cancer; gastrointestinal system cancer (multiple); and lung cancer (multiple). Orthologous to human FOXA1 (forkhead box A1).

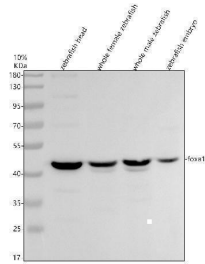
Overview

Product Name	Anti-Zebrafish FOXA1 Antibody Picoband®
Reactive Species	Zebrafish
Description	Boster Bio Anti-Zebrafish FOXA1 Antibody Picoband® catalog # AZQ6POC9. Tested in WB, IF 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	IF, 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	Q6POC9

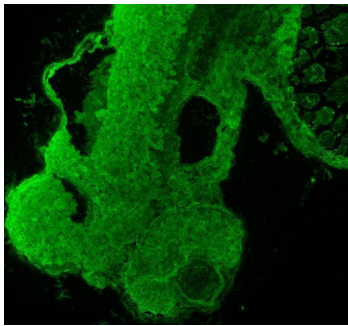
Technical Details

Immunogen	E.coli-derived Zebrafish FOXA1 recombinant protein (Position: M1-Q416).
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 Immunofluorescence, 2 ug/ml, Zebrafish

Anti-Zebrafish FOXA1 Antibody Picoband® (AZQ6P0C9) Images



Western blot analysis of FOXA1 using anti-FOXA1 antibody (AZQ6P0C9). Electrophoresis was performed on a 10% 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, Lane 4: zebrafish embryo 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-FOXA1 antigen affinity purified polyclonal antibody (AZQ6P0C9) 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 FOXA1 at approximately 46 kDa. The expected band size for FOXA1 is at 46 kDa.



IF analysis of FOXA1 using anti-FOXA1 antibody (AZQ6P0C9). FOXA1 was detected in a paraffin-embedded section of zebrafish embryo tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH 8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 5 ug/mL rabbit anti-FOXA1 Antibody (AZQ6P0C9) overnight at 4°C. DyLight®488 Conjugated Goat Anti-Rabbit IgG (BA1127) was used as secondary antibody at 1:500 dilution and incubated for 30 minutes at 37°C. The section was counterstained with DAPI. Visualize using a fluorescence microscope and filter sets appropriate for the label used.

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