

Anti-Zebrafish Vimentin/VIM Antibody Picoband®

Catalog Number: AZQ9YHX5

About VIM

Predicted to be a structural constituent of cytoskeleton. Acts upstream of or within wound healing. Predicted to be active in several cellular components, including axon; intermediate filament; and plasma membrane. Is expressed in several structures, including axis; integument; midbrain neural keel; nervous system; and neural tube. Human ortholog(s) of this gene implicated in atherosclerosis; autoimmune disease (multiple); and cataract 30. Orthologous to human VIM (vimentin).

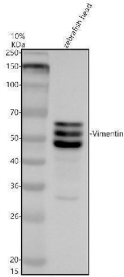
Overview

Product Name	Anti-Zebrafish Vimentin/VIM Antibody Picoband®
Reactive Species	Zebrafish
Description	Boster Bio Anti-Zebrafish Vimentin/VIM Antibody Picoband® catalog # AZQ9YHX5. Tested in WB, IHC 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	IHC, 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	Q9YHX5

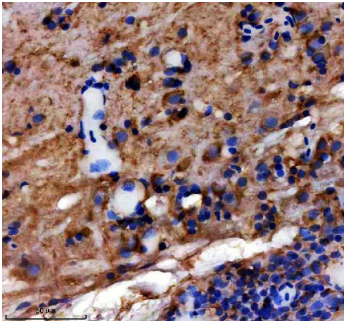
Technical Details

Immunogen	E.coli-derived Zebrafish Vimentin/VIM recombinant protein (Position: S63-N362).
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 Immunohistochemistry(Paraffin-embedded Section), 2-5 ug/ml, Zebrafish

Anti-Zebrafish Vimentin/VIM Antibody Picoband® (AZQ9YHX5) Images



Western blot analysis of Vimentin/VIM using anti-Vimentin/VIM antibody (AZQ9YHX5). 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. 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-Vimentin/VIM antigen affinity purified polyclonal antibody (AZQ9YHX5) 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 Vimentin/VIM at approximately 54 kDa. The expected band size for Vimentin/VIM is at 54 kDa.



IHC analysis of Vimentin/VIM using anti-Vimentin/VIM antibody (AZQ9YHX5). Vimentin/VIM was detected in a paraffin-embedded section of zebrafish brain 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 2 ug/ml rabbit anti-Vimentin/VIM Antibody (AZQ9YHX5) overnight at 4°C. Peroxidase Conjugated Goat Anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit (Catalog # SV0002) with DAB as the chromogen.

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