

Anti-Zebrafish GPRC5BA Antibody Picoband®

Catalog Number: AZX1WEL2

About GPRC5BA

Predicted to enable protein kinase activator activity and protein kinase binding activity. Acts upstream of or within lymph vessel development. Predicted to be located in membrane. Predicted to be part of receptor complex. Predicted to be active in extracellular exosome; intracellular membrane-bounded organelle; and plasma membrane. Human ortholog(s) of this gene implicated in megalencephalic leukoencephalopathy with subcortical cysts. Orthologous to human GPRC5B (G protein-coupled receptor class C group 5 member B).

Submit a product review to Biocompare.com

[Submit a review of this product](#) to Biocompare.com to receive a \$20 Amazon.com giftcard! Your reviews help your fellow scientists make the right decisions. Thank you for your contribution.



Overview

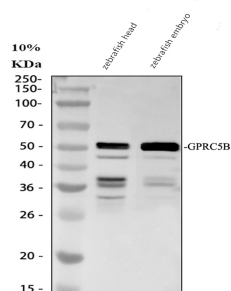
Product Name	Anti-Zebrafish GPRC5BA Antibody Picoband®
Reactive Species	Zebrafish
Description	Boster Bio Anti-Zebrafish GPRC5BA Antibody Picoband® catalog # AZX1WEL2. Tested in WB, IHC applications. This antibody reacts with Zebrafish. The brand Picoband indicates this is a premium antibody that guarantees 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	X1WEL2

Technical Details

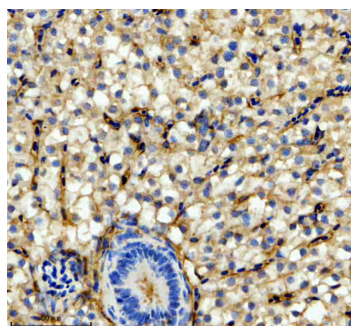
Immunogen	E.coli-derived Zebrafish GPRC5BA recombinant protein (Position: E290-W397).
Form	Lyophilized

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

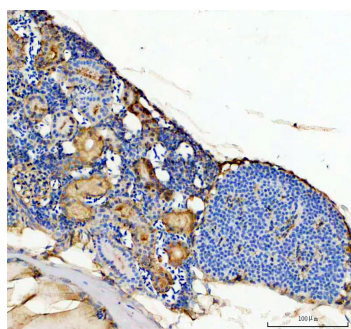
Anti-Zebrafish GPRC5BA Antibody Picoband® (AZX1WEL2) Images



Western blot analysis of GPRC5BA using anti-GPRC5BA antibody (AZX1WEL2). 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: 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-GPRC5BA antigen affinity purified polyclonal antibody (AZX1WEL2) 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 GPRC5BA at approximately 50 kDa. The expected band size for GPRC5BA is at 42 kDa.



IHC analysis of GPRC5BA using anti-GPRC5BA antibody (AZX1WEL2). GPRC5BA was detected in a paraffin-embedded section of zebrafish liver 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-GPRC5BA Antibody (AZX1WEL2) 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.



IHC analysis of GPRC5BA using anti-GPRC5BA antibody (AZX1WEL2). GPRC5BA was detected in a paraffin-embedded section of zebrafish kidney 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-GPRC5BA Antibody (AZX1WEL2) 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.

[Anti-Zebrafish GPRC5BA Antibody](#)

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