

## Anti-Zebrafish ak2 Antibody (Center)

Catalog Number: M04253

### About ak2

Catalyzes the reversible transfer of the terminal phosphate group between ATP and AMP. Plays an important role in cellular energy homeostasis and in adenine nucleotide metabolism. Adenylate kinase activity is critical for regulation of the phosphate utilization and the AMP de novo biosynthesis pathways. Plays a key role in hematopoiesis.

### Overview

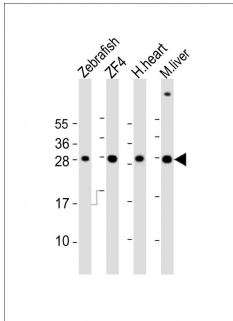
Product Name	Anti-Zebrafish ak2 Antibody (Center)
Reactive Species	Human, Mouse, Zebrafish
Description	Boster Bio Anti-Zebrafish ak2 Antibody (Center) (Catalog # M04253). Tested in WB, IHC-P, Flow Cytometry application(s). This antibody reacts with Human, Mouse, Zebrafish.
Application	Flow Cytometry, IHC-P, WB
Clonality	Polyclonal
Formulation	Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide.
Storage Instructions	Maintain refrigerated at 2-8°C for up to 2 weeks. For long-term storage, store at -20°C in small aliquots to prevent freeze-thaw cycles.
Host	Rabbit
Uniprot ID	Q1L8L9

### Technical Details

Immunogen	This Zebrafish ak2 antibody is generated from a rabbit immunized with a KLH conjugated synthetic peptide between 147-183 amino acids from the Central region of Zebrafish ak2.
Predicted Reactive Species	Chicken, Mouse
Isotype	Rabbit IgG
Purification	This antibody is purified through a protein A column, followed by peptide affinity purification.
Suggested Dilutions	Dilute the sample so that the expected range of concentrations fall within the detection range of this kit. If the expected range of concentration is unknown, a pilot test should be conducted to decide the optimal dilution ratio for your samples. Some PubMed article(s) citing the expression level of this target are as follows: Boster Bio's internal QC testing used: WB: 1:2000 IHC-P: 1:25

	FC: 1:25
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## Anti-Zebrafish ak2 Antibody (Center) (M04253) Images



All lanes : Anti-Zebrafish ak2 Antibody (Center) at 1:2000 dilution

Lane 1: Zebrafish lysates

Lane 2: ZF4 whole cell lysates

Lane 3: human heart lysates

Lane 4: mouse liver lysates

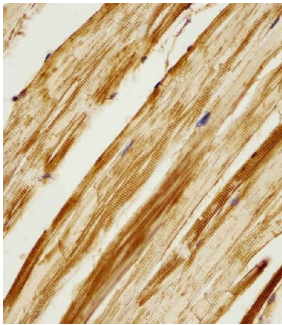
Lysates/proteins at 20 µg per lane.

Secondary

Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution.

Predicted band size : 27 kDa

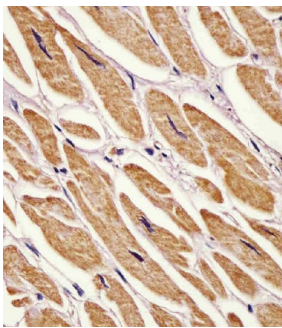
Blocking/Dilution buffer: 5% NFDM/TBST.



M04253 staining Zebrafish ak2 in zebra fish body tissue sections by Immunohistochemistry (IHC-P

-paraformaldehyde-fixed, paraffin-embedded sections).

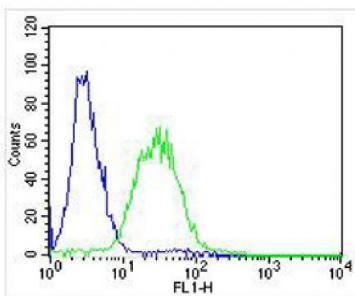
Tissue was fixed with formaldehyde and blocked with 3% BSA for 0.5 hour at room temperature; antigen retrieval was by heat mediation with a citrate buffer (pH6). Samples were incubated with primary antibody (1/25) for 1 hour at 37°C. A undiluted biotinylated goat polyvalent antibody was used as the secondary antibody.



M04253 staining Zebrafish ak2 in human heart tissue sections by Immunohistochemistry (IHC-P

-paraformaldehyde-fixed, paraffin-embedded sections).

Tissue was fixed with formaldehyde and blocked with 3% BSA for 0.5 hour at room temperature; antigen retrieval was by heat mediation with a citrate buffer (pH6). Samples were incubated with primary antibody (1/25) for 1 hour at 37°C. A undiluted biotinylated goat polyvalent antibody was used as the secondary antibody.



Overlay histogram showing ZF4 cells stained with M04253 (green line). The cells were fixed with 2% paraformaldehyde (10 min) and then permeabilized with 90% methanol for 10 min. The cells were then incubated in 2% bovine serum albumin to block non-specific protein-protein interactions followed by the antibody (M04253, 1:25 dilution) for 60 min at 37°C. The secondary antibody used was Goat-Anti-Rabbit IgG, DyLight® 488 Conjugated Highly Cross-Adsorbed at 1/400 dilution for 40 min at 37°C. Isotype control antibody (blue line) was rabbit IgG (1g/1x10<sup>6</sup> cells) used under the same conditions. Acquisition of >10,000 events was performed.

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