

## Anti-LRRC4 Antibody Picoband®

Catalog Number: A07447-1

### About LRRC4

Predicted to be involved in modulation of chemical synaptic transmission and synapse organization. Predicted to act upstream of or within synapse organization. Predicted to be located in dendritic spine; excitatory synapse; and postsynaptic membrane. Predicted to be active in Schaffer collateral - CA1 synapse; glutamatergic synapse; and postsynaptic density membrane.

### Overview

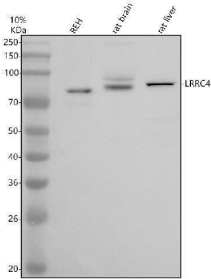
Product Name	Anti-LRRC4 Antibody Picoband®
Reactive Species	Human, Rat
Description	Boster Bio Anti-LRRC4 Antibody Picoband® catalog # A07447-1. Tested in WB, Flow Cytometry, ELISA applications. This antibody reacts with Human, Rat. The brand Picoband indicates this is a premium antibody that guarantees 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	ELISA, Flow Cytometry, WB
Clonality	Polyclonal
Formulation	Each vial contains 4 mg Trehalose, 0.9 mg NaCl, 0.2 mg Na <sub>2</sub> HPO <sub>4</sub> .
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	Q9HBW1

### Technical Details

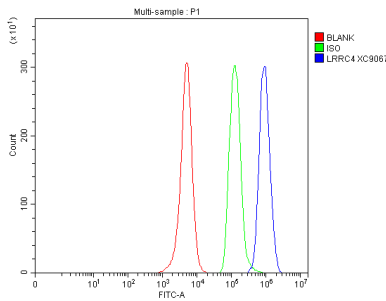
Immunogen	E.coli-derived human LRRC4 recombinant protein (Position: R78-H643).
Form	Lyophilized
Concentration	Adding 0.2 ml of distilled water will yield a concentration of 500 ug/ml.
Purification	Immunogen affinity purified.
Suggested Dilutions	Western blot, 0.25-0.5 ug/ml, Human, Rat Flow Cytometry (Fixed), 1-3 ug/1x10 <sup>5</sup> cells, Human ELISA, 0.1-0.5 ug/ml



## Anti-LRRC4 Antibody Picoband® (A07447-1) Images



Western blot analysis of LRRC4 using anti-LRRC4 antibody (A07447-1). 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: human REH whole cell lysates, Lane 2: rat brain tissue lysates, Lane 3: rat liver 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-LRRC4 antigen affinity purified polyclonal antibody (A07447-1) 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 LRRC4 at approximately 80 kDa. The expected band size for LRRC4 is at 73 kDa.



Flow Cytometry analysis of SH-SY5Y cells using anti-LRRC4 antibody (A07447-1). Overlay histogram showing SH-SY5Y cells stained with A07447-1 (Blue line). The cells were fixed with 4% paraformaldehyde and blocked with 10% normal goat serum. And then incubated with rabbit anti-LRRC4 Antibody (A07447-1, 1 ug/1x10<sup>6</sup> cells) for 30 min at 20°C. Fluoro488 conjugated goat anti-rabbit IgG (BA1127, 5-10 ug/1x10<sup>6</sup> cells) was used as secondary antibody for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG (1 ug/1x10<sup>6</sup>) used under the same conditions. Unlabelled sample without incubation with primary antibody and secondary antibody (Red line) was used as a blank control.

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### Anti-LRRC4 Antibody

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