

## Anti-DENR Antibody

Catalog Number: M06362

### About DENR

May be involved in the translation of target mRNAs by scanning and recognition of the initiation codon. Involved in translation initiation; promotes recruitment of aminoacylated initiator tRNA to P site of 40S ribosomes. Can promote release of deacylated tRNA and mRNA from recycled 40S subunits following ABCE1-mediated dissociation of post-termination ribosomal complexes into subunits. Plays a role in the modulation of the translational profile of a subset of cancer-related mRNAs when recruited to the translational initiation complex by the oncogene MCTS1.

### Overview

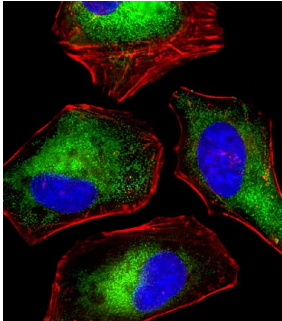
|                      |  |
|----------------------|--|
| Product Name         | Anti-DENR Antibody   |
| Reactive Species     | Human  |
| Description          | Boster Bio Anti-DENR Antibody (Catalog # M06362). Tested in WB, IHC-P, Flow Cytometry, IF application(s). This antibody reacts with Human. |
| Application          | Flow Cytometry, IF, IHC-P, WB  |
| Clonality            | Monoclonal 1542CT106.51.79   |
| Formulation          | Purified monoclonal 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                 | Mouse  |
| Uniprot ID           | O43583   |

### Technical Details

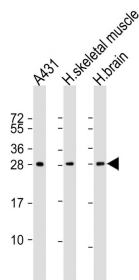
|                            |  |
|----------------------------|--|
| Immunogen                  | This DENR antibody is generated from a mouse immunized with a recombinant protein of human DENR.   |
| Predicted Reactive Species | Human, Mouse   |
| Isotype                    | IgG2b,kappa  |
| Purification               | This antibody is purified through a protein G column, followed by dialysis against PBS.  |
| Suggested Dilutions        | Dilute the sample so that the expected range of concentrations fall within the detection range of this kit.<br>If the expected range of concentration is unknown, a pilot test should be conducted to decide the optimal dilution ratio for your samples.<br>Some PubMed article(s) citing the expression level of this target are as follows:<br>Boster Bio's internal QC testing used:<br>IF: 1:25 |

|  |  |
|--|--|
|  | WB: 1:1000-1:2000<br>IHC-P: 1:25<br>FC: 1:25 |
|--|--|

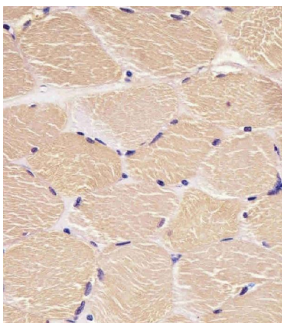
## Anti-DENR Antibody (M06362) Images



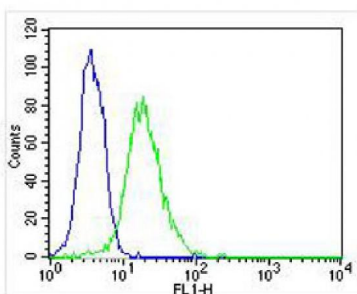
Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa (human cervical epithelial adenocarcinoma cell line) cells labeling DENR with M06362 at 1/25 dilution, followed by Dylight® 488-conjugated goat anti-mouse IgG secondary antibody at 1/200 dilution (green). Immunofluorescence image showing cytoplasm staining on HeLa cell line. Cytoplasmic actin is detected with Dylight® 554 Phalloidin at 1/100 dilution (red). The nuclear counter stain is DAPI (blue).



All lanes : Anti-DENR Antibody at 1:1000-1:2000 dilution  
Lane 1: A431 whole cell lysate  
Lane 2: human skeletal muscle lysate  
Lane 3: human brain lysate  
Lysates/proteins at 20 µg per lane.  
Secondary  
Goat Anti-mouse IgG, (H+L), Peroxidase conjugated at 1/10000 dilution.  
Predicted band size : 22 kDa  
Blocking/Dilution buffer: 5% NFDM/TBST.



M06362 staining DENR in human skeletal muscle 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 hours at 37°C. A undiluted biotinylated goat polyvalent antibody was used as the secondary antibody.



Overlay histogram showing HeLa cells stained with M06362 (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 (M06362, 1:25 dilution) for 60 min at 37°C. The secondary antibody used was Goat-Anti-Mouse IgG, DyLight® 488 Conjugated Highly Cross-Adsorbed at 1/400 dilution for 40 min at 37°C. Isotype control antibody (blue line) was mouse IgG2b (1g/1x10<sup>6</sup> cells) used under the same conditions. Acquisition of >10,000 events was performed.

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