

# Anti-Human ESRRB, monoclonal

**Alternate Names:** Steroid hormone receptor ERR2

**Cat. No.** 15-024

FOR RESEARCH USE ONLY

NOT FOR USE IN HUMANS



## PRODUCT DESCRIPTION

### UniProt Summary

#### UniProt

Primary Accession: [O95718](#)

Nuclear receptor, may regulate ESR1 transcriptional activity. Induces the expression of PERM1 in the skeletal muscle. Binds DNA as a monomer. Nucleus. Event=Alternative splicing; Named isoforms=3; Name=1; Synonyms=ERRbeta2-delta10; Isold=O95718-1; Sequence=Displayed; Name=2; Isold=O95718-2; Sequence=VSP\_042211; Name=3; Synonyms=ERRbeta-short-form; Isold=O95718-3; Sequence=VSP\_042212; Acetylated by PCAF/KAT2 (in vitro). Deafness, autosomal recessive, 35 (DFNB35) [MIM:608565]: A form of non-syndromic deafness characterized by non-progressive, prelingual hearing loss. Note=The disease is caused by mutations affecting the gene represented in this entry. Belongs to the nuclear hormone receptor...

### Physical Characteristics

**Quantity:** 1 ml (culture supernatant) or 100ug at 1mg/ml (purified)

**Format:** culture supernatant or purified material

**Host/Isotype:** mouse IgG1

**Clonality:** monoclonal; ID R1026.1.2A6

**Formulation:** culture supernatant contains 0.02% NaN<sub>3</sub>. Purified material contains 30% glycerol, PBS and 0.02% NaN<sub>3</sub>

**Specificity:** monospecific for human ESRRB ; see "Microarray Analysis" below

**Reactivity:** human; not tested for cross reactivity in other species

**Stability/Storage:** 12 months long term: -20°C; short term: 4°C; avoid freeze-thaw cycles; aliquot as required

**Handling Notes:** small volumes of antibody may occasionally become entrapped in the seal of the product vial during shipment and storage; briefly centrifuge the vial on a tabletop centrifuge to dislodge any liquid in the container cap

### Tested Research Applications

- Western Immunoblotting - Not recommended
- Immunoprecipitation - PASS by SOP

Antibody tested as purified IgG. Optimal dilution to be determined by user.

### Quality Assurance

#### Notes:

1. Please refer to the SOP manual ([click here](#)) to for detailed explanation of the experimental setup and thresholds used to evaluate the results shown below.
2. Predicted MW of the target protein expressed in the western immunoblotting (WB) and immprecipitation (IP) experiment is **55.0kDa** and the three fusion tags (venus, 3xFLAG, V5) adds another **38.7kDa** to this protein. Therefore the protein is expected to migrate **~93.7kDa** in a denatured SDS-PAGE gel.
3. See results below for any applicable cautionary notes.

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Lot-specific COA version tracker: v1.0.0



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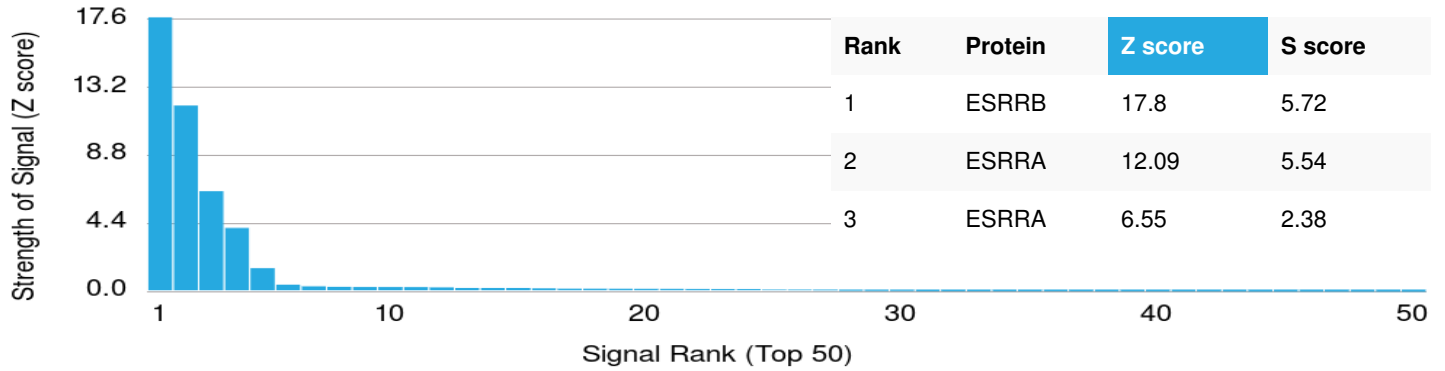
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NEXTGEN PROTEOMICS

## PRODUCT DESCRIPTION

### Quality Assurance (HuProt™ Array)



**About Z and S Scores:** The z-score represents the strength of a signal that a mAb (in combination with a fluorescently-tagged anti-IgG secondary antibody) produces when binding to a particular protein on the HuProt™ array. Z-scores are described in units of standard deviations (SDs) above the mean value of all signals generated on that array. To be considered a definite binding event between a mAb and a protein target, the z-score must have a value of at least 2.5 (SDs above the mean). The s-score represents the relative target specificity of a mAb, and is the difference (also in units of SDs) between the z-score a mAb generates upon binding to one protein and the next highest z-score that the mAb generates against another protein. For example, if a mAb binds to protein X with an z-score of 43 and to protein Y with an z-score of 14, then the s-score for the binding of that mAb to protein X is equal to 29.

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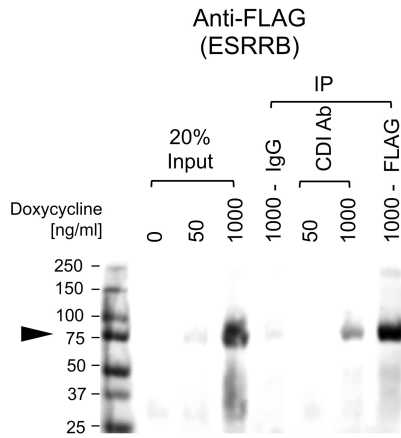


NEXTGEN PROTEOMICS

PRODUCT DESCRIPTION

## Quality Assurance (continued)

### IMMUNOPRECIPITATION - Predicted MW 93.7kDa



Tet-ON HeLa cells were transfected with construct encoding ESRRB (BC132595) with an N-terminal fusion of FLAG, YFP (Venus) and V5 tags under a tet-inducible promoter. These cells were stimulated with 0, 50 or 1000 ng/ml doxycycline. Immunoprecipitation (IP) was carried out using 5 µg of either IgG, CDI mAb ESRRB (cloneID# R1026.1.2A6) or 1 µg of FLAG-M2. Immunoblotting was performed using rabbit Anti-FLAG (1:1000, Cell Signaling #2368).

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