Applications

WB,ELISA



TRβ1 (Phospho Ser142) Rabbit pAb

CatalogNo: YP0479 Orthogonal Validated 💽

Key Features

Host Species Reactivity

Rabbit
 Human, Mouse, Rat

MW Isotype
• 45kD (Observed) • IgG

Recommended Dilution Ratios

WB 1:500-1:2000 ELISA 1:5000

Not yet tested in other applications.

Storage

Storage* -15°C to -25°C/1 year(Do not lower than -25°C)

Formulation Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Basic Information

Clonality Polyclonal

Immunogen Information

Immunogen The antiserum was produced against synthesized peptide derived from human TR-beta1

around the phosphorylation site of Ser142. AA range:116-165

Specificity Phospho-TRβ1 (S142) Polyclonal Antibody detects endogenous levels of TRβ1 protein

only when phosphorylated at S142. The name of modified sites may be influenced by many factors, such as species (the modified site was not originally found in human samples) and the change of protein sequence (the previous protein sequence is incomplete, and the protein sequence may be prolonged with the development of protein sequencing technology). When naming, we will use the "numbers" in historical reference to keep the sites consistent with the reports. The antibody binds to the following modification sequence (lowercase letters are modification sites):HPsYS

Target Information

Gene name

THRB

Protein Name

Thyroid hormone receptor beta

Organism	Gene ID	UniProt ID
Human	<u>7068;</u>	<u>P10828;</u>
Mouse	<u>21834;</u>	<u>P37242;</u>
Rat		<u>P18113;</u>

Cellular Localization

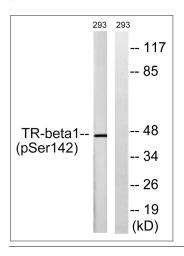
Nucleus.

Tissue specificity Brain, Kidney, Pituitary, Placenta, Testis,

Function

Disease: Defects in THRB are the cause of generalized thyroid hormone resistance (GTHR) [MIM:188570, 274300]. GTHR is transmitted as an autosomal dominant trait, but an autosomal recessive form also exists. The disease is characterized by goiter, abnormal mental functions, increased susceptibility to infections, abnormal growth and bone maturation, tachycardia and deafness. Affected individuals may also have attention deficithyperactivity disorders (ADHD) and language difficulties. GTHR patients also have high levels of circulating thyroid hormones (T3-T4), with normal or slightly elevated thyroid stimulating hormone (TSH)., Disease: Defects in THRB are the cause of selective pituitary thyroid hormone resistance (PRTH) [MIM:145650]; also called familial hyperthyroidism due to inappropriate thyrotropin secretion. PRTH is a variant form of thyroid hormone resistance and is characterized by clinical hyperthyroidism, with elevated free thyroid hormones, but inappropriately normal serum TSH. Unlike GRTH, where the syndrome usually segregates with a dominant allele, the mode of inheritance in PRTH has not been established., Domain: Composed of three domains: a modulating N-terminal domain, a DNAbinding domain and a C-terminal steroid-binding domain., Function: High affinity receptor for triiodothyronine., similarity: Belongs to the nuclear hormone receptor family. NR1 subfamily., similarity: Contains 1 nuclear receptor DNA-binding domain., subunit: Interacts with NOCA7 in a ligand-inducible manner. Interacts with C1D.,

Validation Data



Western blot analysis of lysates from 293 cells treated with PMA 125ng/ml 30', using TR-beta1 (Phospho-Ser142) Antibody. The lane on the right is blocked with the phospho peptide.

| Contact information

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Please scan the QR code to access additional product information: TRβ1 (Phospho Ser142) Rabbit pAb

For Research Use Only. Not for Use in Diagnostic Procedures.

Antibody | ELISA Kits | Protein | Reagents