## RHEBL1 Antibody

Catalog No: #42968



Orders: order@signalwayantibody.com Support: tech@signalwayantibody.com

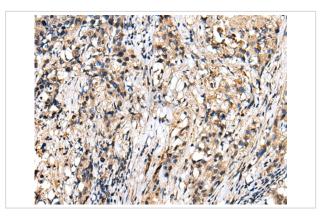
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| Product Name          | RHEBL1 Antibody   |
|-----------------------|---|
| Host Species          | Rabbit  |
| Clonality             | Polyclonal  |
| Purification          | Antigen affinity purification.                                  |
| Applications          | IHC   |
| Species Reactivity    | Hu  |
| Specificity           | The antibody detects endogenous levels of total RHEBL1 protein. |
| Immunogen Type        | protein   |
| Immunogen Description | Fusion protein of human RHEBL1                                  |
| Target Name           | RHEBL1  |
| Other Names           | RHEBL1c   |
| Accession No.         | Swiss-Prot#: Q8TAI7Gene ID: 121268                              |
| Uniprot               | Q8TAI7  |
| GeneID                | 121268;   |
| Concentration         | 0.6mg/ml  |
| Formulation           | Rabbit IgG in pH7.4 PBS, 0.05% NaN3, 40% Glycerol.              |
| Storage               | Store at -20°C  |
|                       |   |

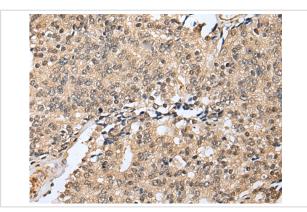
## Application Details

Immunohistochemistry: 1:25-1:100

## **Images**



Immunohistochemical analysis of paraffin-embedded Human lung cancer tissue using #42968 at dilution 1/25.



Immunohistochemical analysis of paraffin-embedded Human prostate cancer tissue using #42968 at dilution 1/25.

## Background

RhebL1 (ras homolog enriched in brain-like protein 1), also known as Rheb2 or GTPase RhebL1, is a 183 amino acid protein that belongs to the small GTPase superfamily and Rheb family. Localizing to the cell membrane as well as the cytoplasm, RhebL1 is ubiquitously expressed and is increased two-fold in many tumor cell lines. RhebL1 exhibits GTPase activity and may activate NF-kappa-B-mediated gene transcription. Regulating the activity of Rictor, RhebL1 also promotes signal transduction. RhebL1 exists as two alternatively spliced isoforms and is encoded by a gene that maps to human chromosome 12q13.12 and mouse chromosome 15 F1. Human chromosome 12 encodes over 1,100 genes and comprises approximately 4.5% of the human genome. Chromosome 12 is associated with a variety of diseases and afflictions, including hypochondrogenesis, achondrogenesis, Kniest dysplasia, Noonan syndrome and trisomy 12p, which causes facial developmental defects and seizure disorders. Binds GTP and exhibits intrinsic GTPase activity. May activate NF-kappa-B-mediated gene transcription. Promotes signal transduction through MTOR, activates RPS6KB1, and is a downstream target of the small GTPase-activating proteins TSC1 and TSC2.

Note: This product is for in vitro research use only