

RHEBL1 抗原(重组蛋白)

- 中文名称: RHEBL1 抗原(重组蛋白)
- 英文名称: RHEBL1 Antigen (Recombinant Protein)
- 别 名: Ras homolog enriched in brain like 1; RHEBL1c
- 相关类别: 抗原
- 储 存: 冷冻 (-20℃)

概述

Fusion protein corresponding to N terminal 150 amino acids of human RHEBL1

技术规格

| Full name: | Ras homolog enriched in brain like 1 |
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| Synonyms: | RHEBL1c |
| Swissprot: | Q8TAI7 |
| Gene Accession: | BC027482 |
| Purity: | >85%, as determined by Coomassie blue stained SDS-PAGE |
| Expression system: | Escherichia coli |
| Tags: | His tag C-Terminus, GST tag N-Terminus |
| Background: | RhebL1 (ras homolog enriched in brain-like protein 1), also kno wn as Rheb2 or GTPase RhebL1, is a 183 amino acid protein th at belongs to the small GTPase superfamily and Rheb family. Lo calizing to the cell membrane as well as the cytoplasm, RhebL1 is ubiquitously expressed and is increased two-fold in many tu mor 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 exi sts as two alternatively spliced isoforms and is encoded by a g |



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ene that maps to human chromosome 12q13.12 and mouse chr omosome 15 F1. Human chromosome 12 encodes over 1,100 g enes and comprises approximately 4.5% of the human genome. Chromosome 12 is associated with a variety of diseases and affl ictions, including hypochondrogenesis, achondrogenesis, Kniest dysplasia, Noonan syndrome and trisomy 12p, which causes faci al developmental defects and seizure disorders. Binds GTP and exhibits intrinsic GTPase activity. May activate NF-kappa-B-medi ated gene transcription. Promotes signal transduction through MTOR, activates RPS6KB1, and is a downstream target of the s mall GTPase-activating proteins TSC1 and TSC2.