

RIMKLB Antibody

Catalog No: #42904

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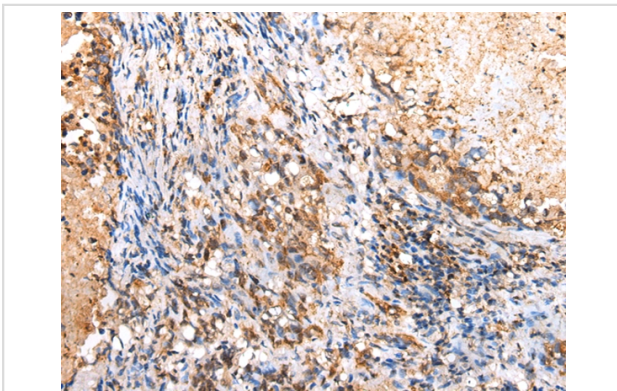
Description

| | |
|-----------------------|---|
| Product Name | RIMKLB Antibody |
| Host Species | Rabbit |
| Clonality | Polyclonal |
| Purification | Antigen affinity purification. |
| Applications | IHC |
| Species Reactivity | Hu |
| Specificity | The antibody detects endogenous levels of total RIMKLB protein. |
| Immunogen Type | protein |
| Immunogen Description | Full length fusion protein of human RIMKLB |
| Target Name | RIMKLB |
| Other Names | NAAGS; FAM80B; NAAGS-I |
| Accession No. | Swiss-Prot#: Q9ULI2 Gene ID: 57494 |
| Uniprot | Q9ULI2 |
| GeneID | 57494; |
| Concentration | 1.6mg/ml |
| Formulation | Rabbit IgG in pH7.4 PBS, 0.05% NaN ₃ , 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 #42904 at dilution 1/30,

Background

RIMKLB (ribosomal modification protein rimK-like family member B), also known as NAAGS (N-acetyl-aspartyl-glutamate synthetase B), beta-citryl-glutamate synthase B or FAM80B, is a 386 amino acid cytoplasmic protein that belongs to the rimK family. Acting as the catalyst in the synthesis of beta-citryl-glutamate and N-acetyl-aspartyl-glutamate, RIMKLB contains one ATP-grasp domain and exists as two alternatively spliced isoforms. The gene encoding RIMKLB maps to human chromosome 12, which 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.

Note: This product is for in vitro research use only