EVA1A Antibody

Catalog No: #40386

Description



Orders: order@signalwayantibody.com

Support: tech@signalwayantibody.com

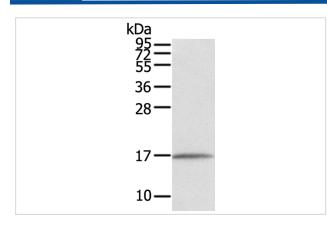
| Product Name          | EVA1A Antibody  |
|-----------------------|---|
| Host Species          | Rabbit  |
| Clonality             | Polyclonal  |
| Purification          | Antigen affinity purification.  |
| Applications          | WB IHC  |
| Species Reactivity    | Hu Ms   |
| Specificity           | The antibody detects endogenous levels of total EVA1A protein.  |
| Immunogen Type        | Peptide   |
| Immunogen Description | Synthetic peptide corresponding to residues near the C terminal of human eva-1 homolog A (C. elegans) |
| Target Name           | EVA1A   |
| Other Names           | FAM176A; TMEM166  |
| Accession No.         | Swiss-Prot:Q9H8M9Gene Accssion:NP_115557  |
| Uniprot               | Q9H8M9  |
| GeneID                | 84141;  |
| SDS-PAGE MW           | 17KD  |
| Concentration         | 2.1mg/ml  |
| Formulation           | Rabbit IgG in pH7.4 PBS, 0.05% NaN3, 40% Glycerol.  |
| Storage               | Store at -20°C  |
|                       |   |

## **Application Details**

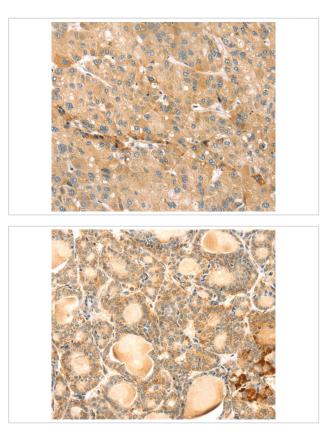
Western blotting: 1:200-1:1000

Immunohistochemistry:1:25-1:100

## Images



Gel: 12%SDS-PAGE Lysate: 40ug Human normal liver tissuePrimary antibody: 1/550 dilution Secondary antibody dilution: 1/8000 Exposure time: 10 seconds



Immunohistochemical analysis of paraffin-embedded Human liver cancer tissue using #40386 at dilution 1/30.

Immunohistochemical analysis of paraffin-embedded Human thyroid cancer tissue using #40386 at dilution 1/30.

## Background

TMEM166, also known as FAM176A (family with sequence similarity 176, member A), is a 152 amino acid protein encoded by a gene mapping to human chromosome 2. The second largest human chromosome, 2 consists of 237 million bases encoding over 1,400 genes and making up approximately 8% of the human genome. A number of genetic diseases are linked to genes on chromosome 2. Harlequin icthyosis, a rare and morbid skin deformity, is associated with mutations in the ABCA12 gene.?

Note: This product is for in vitro research use only