

Anti-TACSTD2 Antibody

Rabbit polyclonal antibody to TACSTD2 Catalog # AP60401

Product Information

Application WB, IF/IC, IHC

Primary Accession P09758
Other Accession Q8BGV3

Reactivity Human, Mouse, Rat, Pig

HostRabbitClonalityPolyclonalCalculated MW35709

Additional Information

Gene ID 4070

Other Names GA733-1; M1S1; TROP2; Tumor-associated calcium signal transducer 2; Cell

surface glycoprotein Trop-2; Membrane component chromosome 1 surface

marker 1; Pancreatic carcinoma marker protein GA733-1

Target/Specificity Recognizes endogenous levels of TACSTD2 protein.

Dilution WB~~WB (1/500 - 1/1000), IHC (1/100 - 1/200), IF/IC (1/100 - 1/500)

IF/IC~~N/A IHC~~WB (1/500 - 1/1000), IHC (1/100 - 1/200), IF/IC (1/100 -

1/500)

Format Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30%

glycerol, and 0.09% (W/V) sodium azide.

Storage Store at -20 °C.Stable for 12 months from date of receipt

Protein Information

Name TACSTD2

Synonyms GA733-1, M1S1, TROP2

Function May function as a growth factor receptor.

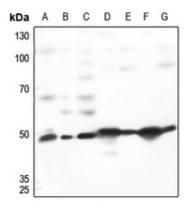
Cellular Location Membrane; Single-pass type I membrane protein.

Tissue Location Placenta, pancreatic carcinoma cell lines.

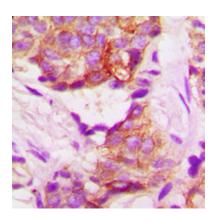
Background

KLH-conjugated synthetic peptide encompassing a sequence within the center region of human TACSTD2. The exact sequence is proprietary.

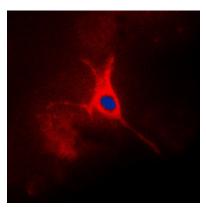
Images



Western blot analysis of TACSTD2 expression in HEK293T (A), Hela (B), H1688 (C), mouse kidney (D), mouse lung (E), rat kidney (F), rat lung (G) whole cell lysates.



Immunohistochemical analysis of TACSTD2 staining in human lung cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of TACSTD2 staining in A549 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.