

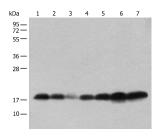
Tel:240-252-7368(USA) Fax: 240-252-7376(USA) techsupport@elabscience.com Website: www.elabscience.com

BUD31 Polyclonal Antibody

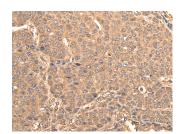
Catalog No.E-AB-18552ReactivityH,M,RStorageStore at -20°C. Avoid freeze / thaw cycles.HostRabbitApplicationsWB,IHC,ELISAIsotypeIgG

Note: Centrifuge before opening to ensure complete recovery of vial contents.

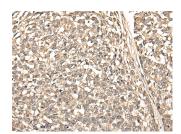
Images



Western Blot analysis of Human testis? Mouse brain Mouse heart? PC-3 cell,231 cell? Raji and Jurkat cell lysates Mouse lung and Rat spleen using BUD31 Polyclonal Antibody at dilution of 1:800.



Immunohistochemistry of paraffinembedded Human liver cancer tissue using BUD31 Polyclonal Antibody at dilution of 1:75(×200)



Immunohistochemistry of paraffinembedded Human esophagus cancer tissue using BUD31 Polyclonal Antibody at dilution of 1:75(×200)

Immunogen Information

Immunogen Full length fusion protein

Gene Accession BC022821 **Swissprot** P41223

Synonyms Bud31,BUD31,Cwc14,EDG 2,EDG2,fSAP17,Protein

EDG-2,Protein G10 homolog,YCR063W

Product Information

Calculated MW 17 kDa

Observed MW Refer to figures

Buffer PBS with 0.05% NaN3 and 40% Glycerol,pH7.4

Purify Antigen affinity purification

Dilution WB 1:1000-1:5000, IHC 1:50-1:300, ELISA

1:5000-1:10000

Background

BUD31 (Protein G10 homolog, EDG-2) is a 144 amino acid protein encoded by the human gene BUD31. BUD31 is a nuclear protein that belongs to the BUD31 (G10) family. BUD31 is found on chromosome 7 which is about 158 milllion bases long, encodes over 1,000 genes and makes up about 5% of the human genome. Chromosome 7 has been linked to osteogenesis imperfecta, Pendred syndrome, lissencephaly, citrullinemia and Shwachman-Diamond syndrome. The deletion of a portion of the long (q) arm of human chromosome 7 is associated with Williams-Beuren syndrome, a condition characterized by mild mental retardation, an unusual comfort and friendliness with strangers and an elfin appearance. Deletions of portions of the q arm of chromosome 7 are also seen in a number of myeloid disorders including cases of acute myelogenous leukemia and myelodysplasia.

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