

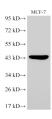
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# **TGFB2 Polyclonal Antibody**

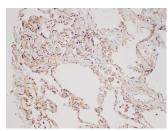
Catalog No.E-AB-40425ReactivityH,M,RStorageStore at -20°C. Avoid freeze / thaw cycles.HostRabbitApplicationsWB,IHCIsotypeIgG

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

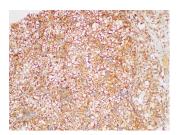
# **Images**



Western Blot analysis of MCF-7 using TGFB2 Polyclonal Antibody at dilution of 1:2000



Immunohistochemistry of paraffinembedded Human lung using TGFB2 Polycloanl Antibody at dilution of 1:200



Immunohistochemistry of paraffinembedded Human clear cell carcinoma of kidney using TGFB2 Polycloanl Antibody at dilution of 1:200

### **Immunogen Information**

Immunogen Recombinant Human Transforming growth factor

beta-2 proprotein protein

**GeneID** 7042 **Swissprot** P61812

**Synonyms** GTSF, LAP, Latency-associated peptide, Polyergin,

TGF-beta2, TGFB2

#### **Product Information**

Calculated MW 47,50 kDa Observed MW 47 kDa

**Buffer** PBS with 0.05% Proclin300, 50% glycerol, pH7.3.

Purify Antigen Affinity Purification

Dilution WB 1:500-1:2000 IHC 1:50-1:200

### **Background**

This gene encodes a member of the transforming growth factor beta (TGFB) family of cytokines, which are multifunctional peptides that regulate proliferation, differentiation, adhesion, migration, and other functions in many cell types by transducing their signal through combinations of transmembrane type I and type II receptors (TGFBR1 and TGFBR2) and their downstream effectors, the SMAD proteins. Disruption of the TGFB/SMAD pathway has been implicated in a variety of human cancers. The encoded protein is secreted and has suppressive effects of interleukin-2 dependent T-cell growth. Translocation t(1;7)(q41;p21) between this gene and HDAC9 is associated with Peters' anomaly, a congenital defect of the anterior chamber of the eye. The knockout mice lacking this gene show perinatal mortality and a wide range of developmental, including cardiac, defects. Alternatively spliced transcript variants encoding different isoforms have been identified.

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