# **Product Name: Treacle Rabbit Polyclonal Antibody**

Catalog #: APRab19231



#### **Summary**

Production Name Treacle Rabbit Polyclonal Antibody

**Description** Rabbit Polyclonal Antibody

**Host** Rabbit

**Application** WB,IHC-P,IF-P,IF-F,ICC/IF,ELISA

**Reactivity** Human, Mouse

### **Performance**

ConjugationUnconjugatedModificationUnmodified

**Isotype** IgG

ClonalityPolyclonalFormLiquid

Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw

cycles.

**Buffer** Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% New type preservative N.

**Purification** Affinity purification

#### **Immunogen**

Storage

Gene Name TCOF1

Alternative Names TCOF1; Treacle protein; Treacher Collins syndrome protein

**Gene ID** 6949.0

Q13428.The antiserum was produced against synthesized peptide derived from human

TCOF1. AA range:41-90

## **Application**

**SwissProt ID** 

WB 1:500-1:2000, IHC-P 1:100-1:300, IF-P/IF-F/ICC/IF 1:200-1:1000, ELISA 1:10000.Not

**Dilution Ratio** 

yet tested in other applications.

Molecular Weight 152kDa

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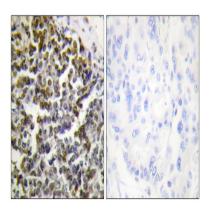


#### **Background**

This gene encodes a nucleolar protein with a LIS1 homology domain. The protein is involved in ribosomal DNA gene transcription through its interaction with upstream binding factor (UBF). Mutations in this gene have been associated with Treacher Collins syndrome, a disorder which includes abnormal craniofacial development. Multiple transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Sep 2008], disease:Defects in TCOF1 are the cause of Treacher Collins syndrome (TCS) [MIM:154500]. TCS is an autosomal dominant disorder of craniofacial development that occurs with an incidence of 1/50,000 live births. The clinical features of TCS are bilaterally symmetrical and include: (1) abnormalities of the external ears, atresia of the external ear canals, and malformation of the middle ear ossicles, which may result in conductive hearing loss; (2) lateral downward sloping of palpebral fissures, frequently with colobomas of the lower eyelids; (3) hypoplasia of the mandible and zygomatic complex; (4) cleft palate.,function:May be involved in nucleolar-cytoplasmic transport. May play a fundamental role in early embryonic development, particularly in development of the craniofacial complex.,PTM:Phosphorylated upon DNA damage, probably by ATM or ATR,,similarity:Contains 1 LisH domain.,

#### **Research Area**

## **Image Data**

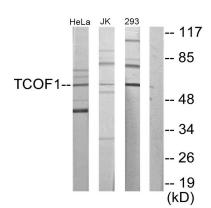


Immunohistochemistry analysis of paraffin-embedded human lung carcinoma tissue, using TCOF1 Antibody. The picture on the right is blocked with the synthesized peptide.

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**C** EnkiLife



Western blot analysis of lysates from Jurkat, 293, HeLa cells, using TCOF1 Antibody. The lane on the right is blocked with the synthesized peptide.

#### Note

For research use only.