



TRI37 rabbit pAb

Catalog No	YP-Ab-08156
Isotype	IgG
Reactivity	Human; Mouse
Applications	WB
Gene Name	TRIM37 KIAA0898 MUL POB1
Immunogen	Synthesized peptide derived from human TRI37 AA range: 315-365
Specificity	This antibody detects endogenous levels of TRI37 at Human/Mouse
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.271% sodium azide.
Source	Polyclonal, Rabbit,IgG
Purification	The antibody was affinity-purified from rabbit serum by affinity-chromatography using specific immunogen.
Dilution	WB 1:500-2000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	E3 ubiquitin protein ligase TRIM37;E3 ubiquitin-protein ligase TRIM37;KIAA898; MUL;MUL protein;Mulibrey nanism gene;Mulibrey nanism protein;POB 1;POB1; RING B box coiled coil protein;TEF 3;TEF3;TRI37_HUMAN;TRIM 37;Trim37; Tripartite motif containing 37;Tripartite motif containing 37 protein;Tripartite motif containing protein 37;Tripartite motif-containing protein 37;TRI37;TRIM37 KIAA0898 MUL POB1;E3 ubiquitin-protein ligase TRIM37 (EC 6.3.2.-) (Mulibrey nanism protein) (Tripartite motif-containing protein 37)
Observed Band	105kD
Calculated Molecular Weight	105kD
Cell Pathway	Cytoplasm, perinuclear region . Peroxisome . Found in vesicles of the peroxisome. Aggregates as aggresomes, a perinuclear region where certain misfolded or aggregated proteins are sequestered for proteasomal degradation. .
Tissue Specificity	Ubiquitous (PubMed:10888877). Highly expressed in testis, while it is weakly expressed in other tissues (PubMed:16310976).
Function	disease:Defects in TRIM37 are the cause of mulibrey nanism (MUL) [MIM:253250]; also called muscle-liver-brain-eye nanism. Mulibrey nanism is an autosomal recessive disorder that involves several tissues of mesodermal origin, implying a defect in a highly pleiotropic gene. Characteristic features include



severe growth failure of prenatal onset and constrictive pericardium with consequent hepatomegaly. In addition, muscle hypotonia, J-shaped sella turcica, yellowish dots in the ocular fundi, typical dysmorphic features and hypoplasia of various endocrine glands causing hormonal deficiency are common., similarity:Belongs to the TRIM/RBCC family.,similarity:Contains 1 B box-type zinc finger.,similarity:Contains 1 MATH domain.,similarity:Contains 1 RING-type zinc finger.,subcellular location:Found in vesicles of the peroxisome.,tissue specificity:Ubiquitous.,

Background

This gene encodes a member of the tripartite motif (TRIM) family, whose members are involved in diverse cellular functions such as developmental patterning and oncogenesis. The TRIM motif includes zinc-binding domains, a RING finger region, a B-box motif and a coiled-coil domain. The RING finger and B-box domains chelate zinc and might be involved in protein-protein and/or protein-nucleic acid interactions. The gene mutations are associated with mulibrey (muscle-liver-brain-eye) nanism, an autosomal recessive disorder that involves several tissues of mesodermal origin. [provided by RefSeq, Mar 2016],

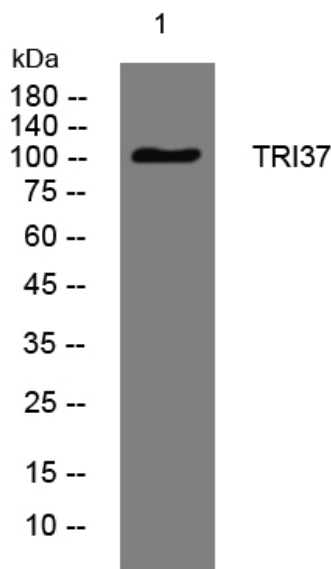
matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images



Western blot analysis of lysates from HpeG2 cells, primary antibody was diluted at 1:1000, 4° over night