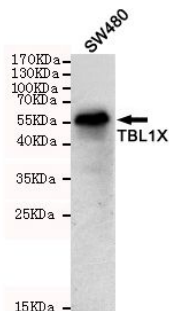




TBL1

Mouse monoclonal Antibody

#53109

**Catalog Number:** 53109**Amount:** 100µg/100µl**Swiss-Prot No. :** O60907**Gene name:** tbl1x**Gene id:** 6907**Clone Number:** 4H2-D5-E9**Form of Antibody:** Purified mouse monoclonal in buffer containing 0.1M Tris-Glycine (pH 7.4, 150 mM NaCl) with 0.2% sodium azide, 50% glycerol**Storage/Stability:** Store at -20°C/1 year**Immunogen:** Purified recombinant human TBL1 protein fragments expressed in E.coli.**Purification:** affinity-chromatography**Specificity/Sensitivity:** This antibody detects endogenous levels of TBL1 and does not cross-react with related proteins**Reactivity:** Human**Applications:** Predicted MW: 58 kd WB: 1:1000 ICC:1:100

Western blot detection of TBL1X in SW480 cell lysates using TBL1X mouse mAb (1:1000 diluted). Predicted band size: 58KDa. Observed band size: 58KDa.

**Background :** The protein encoded by this gene has sequence similarity with members of the WD40 repeat-containing protein family. The WD40 group is a large family of proteins, which appear to have a regulatory function. It is believed that the WD40 repeats mediate protein-protein interactions and members of the family are involved in signal transduction, RNA processing, gene regulation, vesicular trafficking, cytoskeletal assembly and may play a role in the control of cytotypic differentiation. This encoded protein is found as a subunit in corepressor SMRT (silencing mediator for retinoid and thyroid receptors) complex along with histone deacetylase 3 protein. This gene is located adjacent to the ocular albinism gene and it is thought to be involved in the pathogenesis of the ocular albinism with late-onset sensorineural deafness phenotype. Four transcript variants encoding two different isoforms have been found for this gene. This gene is highly similar to the Y chromosome TBL1Y gene.