

## Datasheet

### TBL1X polyclonal antibody (A01)

**Catalog Number:** H00006907-A01

**Regulation Status:** For research use only (RUO)

**Product Description:** Mouse polyclonal antibody raised against a partial recombinant TBL1X.

**Immunogen:** TBL1X (NP\_005638, 478 a.a. ~ 577 a.a) partial recombinant protein with GST tag.

**Sequence:**

LASASFDSTVRLWDIERGVCTHTLTKHQEPVYSVAFSP  
DGKYLASGSFDKCVHIWNTQSGNLVHSYRGTTGGIFEV  
CWNARGDKVGASASDGSVCVLDLRK

**Host:** Mouse

**Reactivity:** Human

**Applications:** ELISA, WB-Ce, WB-Re

(See our web site product page for detailed applications information)

**Protocols:** See our web site at

<http://www.abnova.com/support/protocols.asp> or product page for detailed protocols

**Storage Buffer:** 50 % glycerol

**Storage Instruction:** Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

**Entrez GeneID:** 6907

**Gene Symbol:** TBL1X

**Gene Alias:** EBI, SMAP55, TBL1

**Gene Summary:** 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. [provided by RefSeq]