

## Product datasheet for **CF811698**

### **TBLR1 (TBL1XR1) Mouse Monoclonal Antibody [Clone ID: OTI2A8]**

#### **Product data:**

|                         |  |
|-------------------------|--|
| Product Type:           | Primary Antibodies   |
| Clone Name:             | OTI2A8   |
| Applications:           | WB   |
| Recommended Dilution:   | WB 1:500~2000  |
| Reactivity:             | Human, Rat, Mouse  |
| Host:                   | Mouse  |
| Isotype:                | IgG1   |
| Clonality:              | Monoclonal   |
| Immunogen:              | Full length human recombinant protein of human TBL1XR1 (NP_078941) produced in E.coli.   |
| Formulation:            | Lyophilized powder (original buffer 1X PBS, pH 7.3, 8% trehalose)  |
| Reconstitution Method:  | For reconstitution, we recommend adding 100uL distilled water to a final antibody concentration of about 1 mg/mL. To use this carrier-free antibody for conjugation experiment, we strongly recommend performing another round of desalting process. (OriGene recommends Zeba Spin Desalting Columns, 7KMWCO from Thermo Scientific) |
| Purification:           | Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography (protein A/G)  |
| Conjugation:            | Unconjugated   |
| Storage:                | Store at -20°C as received.  |
| Stability:              | Stable for 12 months from date of receipt.   |
| Predicted Protein Size: | 55.4 kDa   |
| Gene Name:              | TBL1X receptor 1   |
| Database Link:          | <a href="#">NP_078941</a><br><a href="#">Entrez Gene 81004 Mouse</a> <a href="#">Entrez Gene 365755 Rat</a> <a href="#">Entrez Gene 79718 Human</a><br><a href="#">Q9BZK7</a>  |

[View online »](#)

**Background:**

This gene is a member of the WD40 repeat-containing gene family and shares sequence similarity with transducin (beta)-like 1X-linked (TBL1X). The protein encoded by this gene is thought to be a component of both nuclear receptor corepressor (N-CoR) and histone deacetylase 3 (HDAC 3) complexes, and is required for transcriptional activation by a variety of transcription factors. Mutations in these gene have been associated with some autism spectrum disorders, and one finding suggests that haploinsufficiency of this gene may be a cause of intellectual disability with dysmorphism. Mutations in this gene as well as recurrent translocations involving this gene have also been observed in some tumors. [provided by RefSeq, Mar 2016]

**Synonyms:**

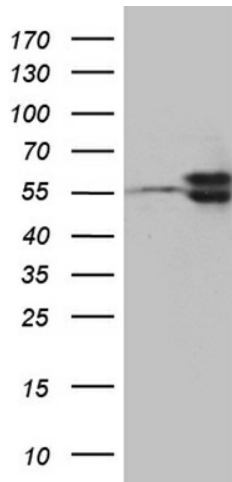
C21; DC42; IRA1; MRD41; TBLR1

**Protein Families:**

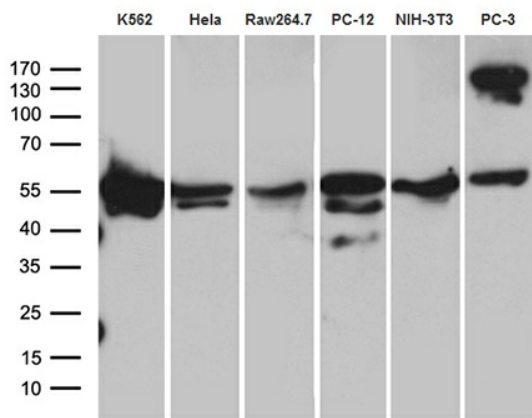
Druggable Genome, Transcription Factors

**Protein Pathways:**

Wnt signaling pathway

**Product images:**


HEK293T cells were transfected with the pCMV6-ENTRY control (Cat# [PS100001], Left lane) or pCMV6-ENTRY TBL1XR1 (Cat# [RC222452], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-TBL1XR1 (Cat# [TA811698])(1:2000). Positive lysates [LY411181] (100ug) and [LC411181] (20ug) can be purchased separately from OriGene.



Western blot analysis of extracts (35ug) from 6 different cell lines by using anti-TBL1XR1 monoclonal antibody (1:500).