

## Product datasheet for **TP504615**

### Sox2 (NM\_011443) Mouse Recombinant Protein

#### Product data:

**Product Type:** Recombinant Proteins

**Description:** Purified recombinant protein of Mouse SRY (sex determining region Y)-box 2 (Sox2), with C-terminal MYC/DDK tag, expressed in HEK293T cells, 20ug

**Species:** Mouse

**Expression Host:** HEK293T

**Expression cDNA Clone or AA Sequence:** >MR204615 protein sequence  
**Red**=Cloning site **Green**=Tags(s)

MYNMMETELKPPGPQQASGGGGGGGNATAAATGGNQKNSPDRVKRPMNAFMVWSRGQRRKMAQENPKMHN  
SEISKRLGAEWKLLSETEKRPFIDEAKRLRALHMKEHPDYKYPKRRKTKLMKKDKYTLPGLLAPGGNS  
MASGVGVGAGLGAGVNRMDSYAHMNGWSNGSYSMMQEQLGYPQHPGLNAHGAAQMMPMHRVDVSALQYN  
SMTSSQTYMNGSPTYSMSYSQQGTPGMALGSMGSSVKSEASSPPVWTSSSHSRAPCQAGDLRDMISMYL  
PGAEVPEPAAPSRLHMAQHYQSGPVPGTAINGLPLSHM

**TRTRPLEQKLISEEDLAANDILDYKDDDDKV**

**Tag:** C-MYC/DDK

**Predicted MW:** 34.4 kDa

**Concentration:** >0.05 µg/µL as determined by microplate BCA method

**Purity:** > 80% as determined by SDS-PAGE and Coomassie blue staining

**Buffer:** 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol

**Note:** For testing in cell culture applications, please filter before use. Note that you may experience some loss of protein during the filtration process.

**Storage:** Store at -80°C after receiving vials.

**Stability:** Stable for 12 months from the date of receipt of the product under proper storage and handling conditions. Avoid repeated freeze-thaw cycles.

**RefSeq:** [NP\\_035573](#)

**Locus ID:** 20674

**UniProt ID:** [P48432](#), [Q60123](#)



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**RefSeq Size:** 2457

**Cytogenetics:** 3 16.93 cM

**RefSeq ORF:** 960

**Synonyms:** lc; lcc; Sox; Sox-2; ys; ysb

**Summary:** This intronless gene encodes a member of the SRY-related HMG-box (SOX) family of transcription factors involved in the regulation of embryonic development and in the determination of cell fate. The product of this gene is required for stem-cell maintenance in the central nervous system, and also regulates gene expression in the stomach. Mutations in a similar gene in human have been associated with optic nerve hypoplasia and with syndromic microphthalmia, a severe form of structural eye malformation. This gene lies within an intron of another gene called SOX2 overlapping transcript (Sox2ot). [provided by RefSeq, Sep 2015]