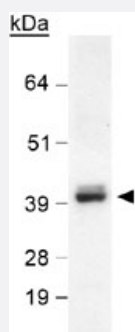


# SOX2 polyclonal antibody

Catalog # PAB12025      Size 100 uL

## Applications



### Western Blot (Tissue lysate)

Western blot analysis of SOX2 in mouse brain lysate with SOX2 polyclonal antibody at 0.5 ug/mL (Cat # PAB12025).

## Specification

<b>Product Description</b>	Rabbit polyclonal antibody raised against synthetic peptide of SOX2.
<b>Immunogen</b>	A synthetic peptide corresponding to amino acids 1-100 of human SOX2.
<b>Host</b>	Rabbit
<b>Reactivity</b>	Chicken, Human, Mouse, Sheep, Xenopus, Zebra fish
<b>Form</b>	Liquid
<b>Quality Control Testing</b>	Antibody Reactive Against Synthetic Peptide.
<b>Recommend Usage</b>	Flow Cytometry (1:100) Immunocytochemistry (1:125-1:250) Immunofluorescence (1:50-1:200) Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) (1:125-1:250) Western Blot (0.5 ug/mL) The optimal working dilution should be determined by the end user.
<b>Storage Buffer</b>	In Tris-glycine, 150 mM NaCl (0.05% sodium azide)
<b>Storage Instruction</b>	Store at -20°C or -80°C. Aliquot to avoid repeated freezing and thawing.

## Note

This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

## Applications

- Western Blot (Tissue lysate)

Western blot analysis of SOX2 in mouse brain lysate with SOX2 polyclonal antibody at 0.5 ug/mL (Cat # PAB12025).

- Immunohistochemistry

## Gene Info — SOX2

Entrez GeneID [6657](#)

Protein Accession# [P48431](#)

Gene Name SOX2

Gene Alias ANOP3, MCOPS3, MGC2413

Gene Description SRY (sex determining region Y)-box 2

Omim ID [184429 206900](#)

Gene Ontology [Hyperlink](#)

**Gene 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 this gene 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 (SOX2 OT). [provided by RefSeq]

**Other Designations** SRY-related HMG-box gene 2|sex-determining region Y-box 2|transcription factor SOX2

## Disease

- [Anophthalmos](#)
- [Coloboma](#)
- [Diabetes Mellitus](#)

- [Diabetic Nephropathies](#)
- [Genetic Predisposition to Disease](#)
- [Kidney Failure](#)
- [Microphthalmos](#)
- [Myopia](#)