

**Product Name: SOX2 Mouse Monoclonal Antibody****Catalog #: AMM80684**

For research use only.

**Summary**

<b>Description</b>	Mouse monoclonal Antibody
<b>Host</b>	Mouse
<b>Application</b>	ICC,ELISA,FC
<b>Reactivity</b>	Human
<b>Conjugation</b>	Unconjugated
<b>Modification</b>	Unmodified
<b>Isotype</b>	Mouse IgG1
<b>Clonality</b>	Monoclonal
<b>Form</b>	Liquid
<b>Concentration</b>	1mg/ml
<b>Storage</b>	Aliquot and store at -20°C (valid for 12 months). Avoid freeze/thaw cycles.
<b>Shipping</b>	Ice bags
<b>Buffer</b>	PBS containing 0.03% sodium azide.
<b>Purification</b>	Affinity Purification

**Application**

<b>Dilution Ratio</b>	ICC 1:50-1:200,ELISA 1:5000-1:20000,FC 1:200-1:400
<b>Molecular Weight</b>	34.3kDa

**Antigen Information**

<b>Gene Name</b>	SOX2
<b>Alternative Names</b>	ANOP3; MCOPS3; MGC2413
<b>Gene ID</b>	6657.0
<b>SwissProt ID</b>	P48431
<b>Immunogen</b>	Purified recombinant fragment of SOX2 (aa1-170) expressed in E. Coli.

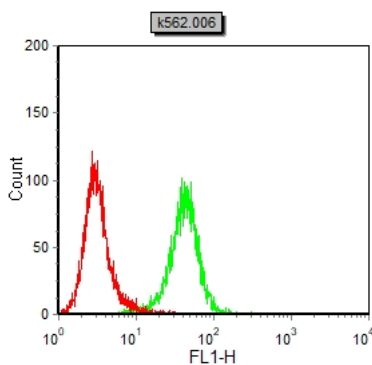
**Background**

SOX2: SRY (sex determining region Y)-box 2. Entrez Protein NP\_003097. It is 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

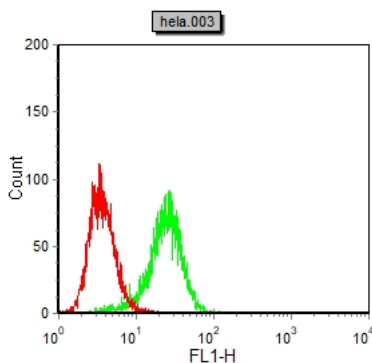
the stomach. Mutations in this gene have been associated with optic nerve hypoplasia and with syndromic microphthalmia, a severe form of structural eye malformation. This gene lies within an intron of another gene called SOX2 overlapping transcript (SOX2OT).

## Research Area

## Image Data



Flow cytometric analysis of K562 cells using SOX2 mouse mAb (green) and negative control (red).



Flow cytometric analysis of HeLa cells using SOX2 mouse mAb (green) and negative control (red).