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## **Datasheet**

## SOX2 purified MaxPab rabbit polyclonal antibody (D01P)

Catalog Number: H00006657-D01P

Regulatory Status: For research use only (RUO)

Product Description: Rabbit polyclonal antibody raised

against a full-length human SOX2 protein.

**Immunogen:** SOX2 (NP\_003097.1, 1 a.a. ~ 317 a.a)

full-length human protein.

## Sequence:

MYNMMETELKPPGPQQTSGGGGGNSTAAAAGGNQK NSPDRVKRPMNAFMVWSRGQRRKMAQENPKMHNSE ISKRLGAEWKLLSETEKRPFIDEAKRLRALHMKEHPDY KYRPRRKTKTLMKKDKYTLPGGLLAPGGNSMASGVG VGAGLGAGVNQRMDSYAHMNGWSNGSYSMMQDQL GYPQHPGLNAHGAAQMQPMHRYDVSALQYNSMTSS QTYMNGSPTYSMSYSQQGTPGMALGSMGSVVKSEA SSSPPVVTSSSHSRAPCQAGDLRDMISMYLPGAEVPE PAAPSRLHMSQHYQSGPVPGTAINGTLPLSHM

Host: Rabbit

Reactivity: Human, Mouse

Applications: WB-Ti

(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: In 1x PBS, pH 7.4

Storage Instruction: Store at -20°C or lower. Aliquot to

avoid repeated freezing and thawing.

Entrez GenelD: 6657

Gene Symbol: SOX2

Gene Alias: ANOP3, MCOPS3, MGC2413

**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 (SOX2OT). [provided by RefSeq]