## SOX2 Conjugated Antibody

Catalog No: #C31128



 Package Size:
 #C31128-AF350 100ul
 #C31128-AF405 100ul
 #C31128-AF488 100ul

 #C31128-AF555 100ul
 #C31128-AF594 100ul
 #C31128-AF647 100ul

 #C31128-AF680 100ul
 #C31128-AF750 100ul
 #C31128-Biotin 100ul

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## Description

Product Name	SOX2 Conjugated Antibody
Host Species	Rabbit
Clonality	Polyclonal
Species Reactivity	Hu
Specificity	The antibody detects endogenous level of total SOX2 protein.
Immunogen Description	Fusion protein corresponding to C terminal 250 amino acids of human SRY (sex determining region Y)-box 2
Conjugates	Biotin AF350 AF405 AF488 AF555 AF594 AF647 AF680 AF750
Other Names	SRY (sex determining region Y)-box 2, ANOP3; MCOPS3
Accession No.	Swiss-Prot#:NCBI Gene ID:NCBI mRNA#:BC013923NCBI Protein#:
Excitation Emission	AF350: 346nm/442nm
	AF405: 401nm/421nm
	AF488: 493nm/519nm
	AF555: 555nm/565nm
	AF594: 591nm/614nm
	AF647: 651nm/667nm
	AF680: 679nm/702nm
	AF750: 749nm/775nm
Calculated MW	34
Formulation	0.01M Sodium Phosphate, 0.25M NaCl, pH 7.6, 5mg/ml Bovine Serum Albumin, 0.02% Sodium Azide
Storage	Store at 4°Cin dark for 6 months

## **Application Details**

Suggested Dilution:	
AF350 conjugated: most applications: 1: 50 - 1: 250	
AF405 conjugated: most applications: 1: 50 - 1: 250	
AF488 conjugated: most applications: 1: 50 - 1: 250	
AF555 conjugated: most applications: 1: 50 - 1: 250	
AF594 conjugated: most applications: 1: 50 - 1: 250	
AF647 conjugated: most applications: 1: 50 - 1: 250	
AF680 conjugated: most applications: 1: 50 - 1: 250	
AF750 conjugated: most applications: 1: 50 - 1: 250	
Biotin conjugated: working with enzyme-conjugated streptavidin, most applications: 1: 50 - 1: 1,000	

Antibodies were produced by immunizing rabbits and were purified by antigen affinity-chromatography.

## Background

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).

Note: This product is for in vitro research use only