

(Tel: 400-999-8863 ■ Emall:Upingbio.163.com



SIX3 Polyclonal Antibody

Catalog No	YP-Ab-05667
Isotype	IgG
Reactivity	Human;Mouse
Applications	WB;ELISA
Gene Name	SIX3
Protein Name	Homeobox protein SIX3 (Sine oculis homeobox homolog 3)
Immunogen	Synthesized peptide derived from part region of human protein
Specificity	SIX3 Polyclonal Antibody detects endogenous levels of protein.
Formulation	Liquid in PBS containing 50% glycerol, and 0.02% sodium azide.
Source	Polyclonal, Rabbit,IgG
Purification	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-2000 ELISA 1:5000-20000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Otorage Glabinty	-20 O/T year
Synonyms	-20 G/T year
-	36kD
Synonyms	
Synonyms Observed Band	36kD
Synonyms Observed Band Cell Pathway	36kD Nucleus .
Synonyms Observed Band Cell Pathway Tissue Specificity	Nucleus . Retina, disease:Defects in SIX3 are the cause of holoprosencephaly type 2 (HPE2) [MIM:157170]. Holoprosencephaly (HPE) [MIM:236100] is the most common structural anomaly of the brain, in which the developing forebrain fails to correctly separate into right and left hemispheres. Holoprosencephaly is genetically heterogeneous and associated with several distinct facies and phenotypic variability.,function:May be involved in visual system development.,similarity:Belongs to the SIX/Sine oculis homeobox
Synonyms Observed Band Cell Pathway Tissue Specificity Function	Nucleus . Retina, disease:Defects in SIX3 are the cause of holoprosencephaly type 2 (HPE2) [MIM:157170]. Holoprosencephaly (HPE) [MIM:236100] is the most common structural anomaly of the brain, in which the developing forebrain fails to correctly separate into right and left hemispheres. Holoprosencephaly is genetically heterogeneous and associated with several distinct facies and phenotypic variability.,function:May be involved in visual system development.,similarity:Belongs to the SIX/Sine oculis homeobox family.,similarity:Contains 1 homeobox DNA-binding domain., This gene encodes a member of the sine oculis homeobox transcription factor family. The encoded protein plays a role in eye development. Mutations in this gene have been associated with holoprosencephaly type 2. [provided by RefSeq,



UpingBio technology Co.,Ltd

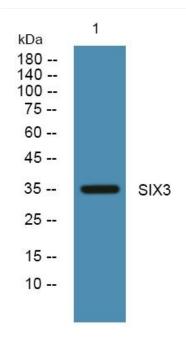




Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images



Western blot analysis of lysates from KB cells, primary antibody was diluted at 1:1000, 4° over night