

## Six1 Polyclonal Antibody

<b>Catalog No :</b>	YT4305
<b>Reactivity :</b>	Human;Mouse
<b>Applications :</b>	WB;ELISA;IHC
<b>Target :</b>	Six1
<b>Fields :</b>	>>Transcriptional misregulation in cancer
<b>Gene Name :</b>	SIX1
<b>Protein Name :</b>	Homeobox protein SIX1
<b>Human Gene Id :</b>	6495
<b>Human Swiss Prot No :</b>	Q15475
<b>Mouse Gene Id :</b>	20471
<b>Mouse Swiss Prot No :</b>	Q62231
<b>Immunogen :</b>	The antiserum was produced against synthesized peptide derived from human SIX1. AA range:111-160
<b>Specificity :</b>	Six1 Polyclonal Antibody detects endogenous levels of Six1 protein.
<b>Formulation :</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
<b>Source :</b>	Polyclonal, Rabbit,IgG
<b>Dilution :</b>	WB 1:500-2000;IHC 1:50-300; ELISA 2000-20000
<b>Purification :</b>	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
<b>Concentration :</b>	1 mg/ml

**Storage Stability :** -15°C to -25°C/1 year(Do not lower than -25°C)

**Observed Band :** 33kD

**Background :** The protein encoded by this gene is a homeobox protein that is similar to the Drosophila &apos;sine oculis&apos; gene product. This gene is found in a cluster of related genes on chromosome 14 and is thought to be involved in limb development. Defects in this gene are a cause of autosomal dominant deafness type 23 (DFNA23) and branchiootic syndrome type 3 (BOS3). [provided by RefSeq, Jul 2008],

**Function :** disease:Defects in SIX1 are the cause of autosomal dominant deafness type 23 (DFNA23) [MIM:605192].,disease:Defects in SIX1 are the cause of branchiootic syndrome type 3 (BOS3) [MIM:608389]. Urinary tract malformations constitute the most frequent cause of chronic renal failure in the first two decades of life. Branchio-oto-renal syndrome (BOR) is an autosomal dominant developmental disorder of kidney and urinary tract malformations with hearing loss. The major feature of BOR is hearing loss (93% of patients), which can be conductive, sensorineural, or both and varies in age of onset.,function:May be involved in limb tendon and ligament development.,similarity:Belongs to the SIX/Sine oculis homeobox family.,similarity:Contains 1 homeobox DNA-binding domain.,tissue specificity:Specifically expressed in skeletal muscle.,

**Subcellular Location :** Nucleus . Cytoplasm.

**Expression :** Specifically expressed in skeletal muscle.

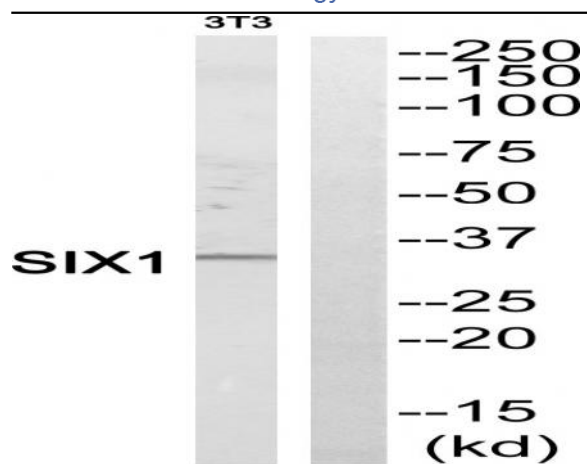
**Sort :** 16350

**No4 :** 1

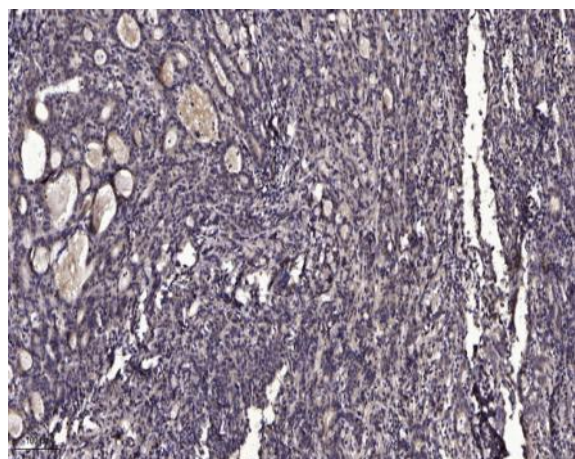
**Host :** Rabbit

**Modifications :** Unmodified

## Products Images



Western blot analysis of SIX1 Antibody. The lane on the right is blocked with the SIX1 peptide.



Immunohistochemical analysis of paraffin-embedded human Gastric adenocarcinoma. 1, Antibody was diluted at 1:200(4° overnight). 2, Tris-EDTA,pH9.0 was used for antigen retrieval. 3,Secondary antibody was diluted at 1:200(room temperature, 45min).