# Six2 (H-4): sc-377193



The Power to Question

## **BACKGROUND**

The Six proteins (sine oculis) are a family of homeodomain transcription factors that share a conserved DNA binding domain. Six2, Six4 (AREC3) and Six5 bind to the same DNA sequence, indicating that they may regulate the same target genes. Six1 and Six4 are both capable of transactivating MEF3 site containing reporter genes, such as myogenin. It has been demonstrated that alterations to homeobox-containing genes may result in cancer. Six1 expression has been shown to be absent or low in normal adult tissues, although it is expressed in several tumor types, including breast carcinoma. Six1 overexpression has been shown to abrogate the  $G_2$  cell cycle checkpoint. Six2 is highly expressed in fetal tissues but expression is limited in adult tissues.

# **CHROMOSOMAL LOCATION**

Genetic locus: SIX2 (human) mapping to 2p21; Six2 (mouse) mapping to 17 E4.

## **SOURCE**

Six2 (H-4) is a mouse monoclonal antibody raised against amino acids 191-230 mapping within an internal region of Six2 of human origin.

#### **PRODUCT**

Each vial contains 200  $\mu$ g IgG<sub>1</sub> kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin. Also available as TransCruz reagent for Gel Supershift and ChIP applications, sc-377193 X, 200  $\mu$ g/0.1 ml.

Six2 (H-4) is available conjugated to agarose (sc-377193 AC), 500  $\mu$ g/0.25 ml agarose in 1 ml, for IP; to HRP (sc-377193 HRP), 200  $\mu$ g/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-377193 PE), fluorescein (sc-377193 FITC), Alexa Fluor® 488 (sc-377193 AF488), Alexa Fluor® 546 (sc-377193 AF546), Alexa Fluor® 594 (sc-377193 AF594) or Alexa Fluor® 647 (sc-377193 AF647), 200  $\mu$ g/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor® 680 (sc-377193 AF680) or Alexa Fluor® 790 (sc-377193 AF790), 200  $\mu$ g/ml, for Near-Infrared (NIR) WB, IF and FCM.

## **APPLICATIONS**

Six2 (H-4) is recommended for detection of Six2 of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2  $\mu$ g per 100-500  $\mu$ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for Six2 siRNA (h): sc-38786, Six2 siRNA (m): sc-38787, Six2 shRNA Plasmid (h): sc-38786-SH, Six2 shRNA Plasmid (m): sc-38787-SH, Six2 shRNA (h) Lentiviral Particles: sc-38786-V and Six2 shRNA (m) Lentiviral Particles: sc-38787-V.

Six2 (H-4) X TransCruz antibody is recommended for Gel Supershift and ChIP applications.

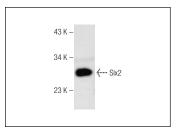
Molecular Weight of Six2: 32 kDa.

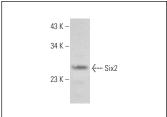
Positive Controls: NIH/3T3 whole cell lysate: sc-2210, Sol8 nuclear extract: sc-2157 or mouse kidney extract: sc-2255.

#### **RECOMMENDED SUPPORT REAGENTS**

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-lgG $\kappa$  BP-HRP: sc-516102 or m-lgG $\kappa$  BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker<sup>TM</sup> Molecular Weight Standards: sc-2035, UltraCruz® Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use m-lgG $\kappa$  BP-FITC: sc-516140 or m-lgG $\kappa$  BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz® Mounting Medium: sc-24941 or UltraCruz® Hard-set Mounting Medium: sc-359850.

## **DATA**





Six2 (H-4): sc-377193. Western blot analysis of Six2 expression in Sol8 nuclear extract.

Six2 (H-4): sc-377193. Western blot analysis of Six2 expression in NIH/3T3 whole cell lysate.

## **SELECT PRODUCT CITATIONS**

- Xu, J., et al. 2014. Tbx18 is essential for normal development of vasculature network and glomerular mesangium in the mammalian kidney. Dev. Biol. 391: 17-31.
- Gao, J., et al. 2020. MES23.5 DA immortalized neuroblastoma cells selfprotect against early injury by overexpressing glial cell-derived neurotrophic factor via Akt1/Eya1/Six2 signaling. J. Mol. Neurosci. 70: 328-339.
- Chang, Y., et al. 2020. MiRNA-516a promotes bladder cancer metastasis by inhibiting MMP9 protein degradation via the Akt/F0X03A/SMURF1 axis. Clin. Transl. Med. 10: e263.

## **STORAGE**

Store at 4° C, \*\*D0 NOT FREEZE\*\*. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

## **RESEARCH USE**

For research use only, not for use in diagnostic procedures.

# **PROTOCOLS**

See our web site at www.scbt.com for detailed protocols and support products.

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