SANTA CRUZ BIOTECHNOLOGY, INC.

HNF-3β (H-4): sc-374376



BACKGROUND

HNF-1 (α and β), HNF-3 (α , β and γ), HNF-4 (α and γ), and HNF-6 compose, in part, a homoeprotein family designated the hepatocyte nuclear factor family. The various HNF-1 isoforms regulate transcription of genes in the liver as well as in other tissues such as kidney, small intestine and thymus. HNF-3 α , HNF-3 β and HNF-3 γ regulate the transcription of numerous hepatocyte genes in adult liver. HNF-3 α and HNF-3 β have also been shown to be involved in gastrulation events such as body axis formation. HNF-4 α and HNF-4 γ have been shown to be important for early embryo development. HNF-4 α is expressed in liver, kidney, pancreas, small intestine, testis and colon; HNF-4 γ is expressed in each of these tissues except liver. HNF-6 has been shown to bind to the promoter of HNF-3 β , which indicates a potential role of HNF-6 in gut endoderm epithelial cell differentiation. Evidence suggests that HNF-6 may also be a transriptional activator for at least 22 other hepatocyte-enriched genes, including cytochrome P450 2C13 and α -1 antitrypsin.

CHROMOSOMAL LOCATION

Genetic locus: FOXA2 (human) mapping to 20p11.21; Foxa2 (mouse) mapping to 2 G3.

SOURCE

HNF-3 β (H-4) is a mouse monoclonal antibody specific for an epitope mapping between amino acids 431-459 at the C-terminus of HNF-3 β of mouse origin.

PRODUCT

Each vial contains 200 μ g lgG₁ 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-374376 X, 200 μ g/0.1 ml.

HNF-3 β (H-4) is available conjugated to agarose (sc-374376 AC), 500 µg/ 0.25 ml agarose in 1 ml, for IP; to HRP (sc-374376 HRP), 200 µg/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-374376 PE), fluorescein (sc-374376 FITC), Alexa Fluor[®] 488 (sc-374376 AF488), Alexa Fluor[®] 546 (sc-374376 AF546), Alexa Fluor[®] 594 (sc-374376 AF594) or Alexa Fluor[®] 647 (sc-374376 AF647), 200 µg/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor[®] 680 (sc-374376 AF680) or Alexa Fluor[®] 790 (sc-374376 AF790), 200 µg/ml, for Near-Infrared (NIR) WB, IF and FCM.

Blocking peptide available for competition studies, sc-374376 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% stabilizer protein).

Alexa Fluor® is a trademark of Molecular Probes, Inc., Oregon, USA

STORAGE

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

PROTOCOLS

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

APPLICATIONS

HNF-3 β (H-4) is recommended for detection of HNF-3 β of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg 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).

HNF-3 β (H-4) is also recommended for detection of HNF-3 β in additional species, including equine, canine, bovine and porcine.

Suitable for use as control antibody for HNF-3 β siRNA (h): sc-35569, HNF-3 β siRNA (m): sc-35570, HNF-3 β shRNA Plasmid (h): sc-35569-SH, HNF-3 β shRNA Plasmid (m): sc-35570-SH, HNF-3 β shRNA (h) Lentiviral Particles: sc-35569-V and HNF-3 β shRNA (m) Lentiviral Particles: sc-35570-V.

HNF-3 β (H-4) X TransCruz antibody is recommended for Gel Supershift and ChIP applications.

Molecular Weight of HNF-3β: 54 kDa.

Positive Controls: Hep G2 cell lysate: sc-2227, MDA-MB-231 cell lysate: sc-2232 or Hep G2 nuclear extract: sc-364819.

DATA





HNF-3 β (H-4) HRP: sc-374376 HRP. Direct western blot analysis of HNF-3 β expression in Hep G2 (**A**) and MDA-MB-231 (**B**) whole cell lysates and Hep G2 nuclear extract (**C**). Cruz Marker^M Molecular Weight Standards detected with Cruz Marker MW Tag-HRP: sc-516732. HNF-3\beta (H-4): sc-374376. Immunofluorescence staining of formalin-fixed Hep G2 cells showing nuclear localization.

SELECT PRODUCT CITATIONS

- Carter, D.A. 2017. Molecular phenotyping of transient postnatal tyrosine hydroxylase neurons in the rat bed nucleus of the stria terminalis. J. Chem. Neuroanat. 82: 29-38.
- An, Y., et al. 2019. Derivation of pluripotent stem cells from nascent undifferentiated teratoma. Dev. Biol. 446: 43-55.
- Malakhova, A.A., et al. 2020. Generation of induced pluripotent stem cell line ICGi018-A from peripheral blood mononuclear cells of a patient with Huntington's disease. Stem Cell Res. 44: 101743.
- Liu, O., et al. 2021. Histone demethylase PHF8 drives neuroendocrine prostate cancer progression by epigenetically upregulating FOXA2.
 J. Pathol. 253: 106-118.

RESEARCH USE

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