SHP (F-12): sc-271469



The Power to Question

BACKGROUND

SHP (short heterodimer partner and small heterodimer partner) is an orphan nuclear receptor containing the dimerization and ligand-binding domains found in other nuclear receptors, but lacking the conserved DNA binding domain. SHP is specifically expressed in liver and other tissues, including fetal liver and adrenal gland, as well as adult spleen and small intestine. In addition, SHP is highly expressed in the murine macrophage cell line RAW 264.7 but suppressed by oxLDL and 13-HODE, which is a ligand for PPARy. SHP interacts with nuclear receptors, including thyroid receptor, retinoic acid receptors (RAR and RXR) and estrogen receptors (ER α and ER β). SHP functions as a negative regulator of these receptors by at least three mechanisms: inhibition of DNA binding via dimerization, direct antagonism of coactivator function through competition and possibly transrepression via recruitment of putative corepressors. In oxLDL-treated, resting macrophage cells, SHP acts as a transcription coactivator of NFkB, suggesting that SHP is a modulatory component in the regulation of the transcriptional activities of NF κ B. Lastly, negative feedback regulation of a hepatic bile acid transporter, NTCP, is controlled by bile acid-activated FXR via induction of SHP to protect the hepatocyte from bile acid-mediated damage in cholestatic conditions.

REFERENCES

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- Lee, H.K., et al. 1998. Structure and expression of the orphan nuclear receptor SHP gene. J. Biol. Chem. 273: 14398-14402.
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- Johansson, L., et al. 2000. The orphan nuclear receptor SHP utilizes conserved LXXLL-related motifs for interactions with ligand-activated estrogen receptors. Mol. Cell. Biol. 20: 1124-1133.
- Klinge, C.M., et al. 2001. Short heterodimer partner (SHP) orphan nuclear receptor inhibits the transcriptional activity of Aryl hydrocarbon receptor (AHR)/AHR nuclear translocator (ARNT). Arch. Biochem. Biophys. 390: 64-70.
- 6. Kim, Y.S., et al. 2001. The orphan nuclear receptor SHP, as a novel coregulator of nuclear factor-κB in oxidized low density lipoprotein-treated macrophage cell RAW 264.7. J. Biol. Chem. 276: 33736-33740.

CHROMOSOMAL LOCATION

Genetic locus: NR0B2 (human) mapping to 1p36.11.

SOURCE

SHP (F-12) is a mouse monoclonal antibody raised against amino acids 1-160 mapping at the N-terminus of SHP of human origin.

PRODUCT

Each vial contains 200 $\mu g \; lg G_{2b}$ kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

APPLICATIONS

SHP (F-12) is recommended for detection of SHP of 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).

Suitable for use as control antibody for SHP siRNA (h): sc-44101, SHP shRNA Plasmid (h): sc-44101-SH and SHP shRNA (h) Lentiviral Particles: sc-44101-V.

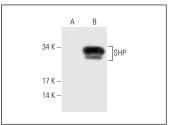
Molecular Weight of SHP: 28 kDa.

Positive Controls: SHP (h): 293T Lysate: sc-114141 or A-431 nuclear extract: sc-2122.

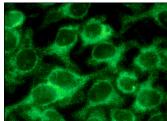
RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-lgG κ BP-HRP: sc-516102 or m-lgG κ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz MarkerTM 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 κ BP-FITC: sc-516140 or m-lgG κ 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



SHP (F-12): sc-271469. Western blot analysis of SHP expression in non-transfected: sc-117752 (**A**) and human SHP transfected: sc-114141 (**B**) 293T whole



SHP (F-12): sc-271469. Immunofluorescence staining of methanol-fixed HeLa cells showing cytoplasmic localization.

SELECT PRODUCT CITATIONS

 Liu, B., et al. 2017. Glutamine attenuates obstructive cholestasis in rats via farnesoid X receptor-mediated regulation of Bsep and Mrp2. Can. J. Physiol. Pharmacol. 95: 215-223.

STORAGE

Store at 4° C, **DO 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.