

FASN Mouse mAb

Catalog Number: bsm-51155M

Target Protein: FASN

Concentration: 1mg/ml

Form: Liquid

Host: Mouse

Clonality: Monoclonal

Clone No.: 2C5

Isotype: IgM

Applications: WB (1:500-5000), IHC-P (1:100-500), IHC-F (1:20-200), IF (1:20-200), ICC/IF (1:100-500)

Reactivity: Human (predicted:Mouse)

Predicted MW: 274 kDa

Entrez Gene: 2194

Swiss Prot: P49327

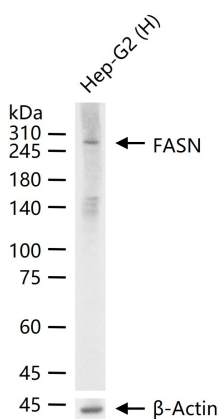
Purification: affinity purified by Protein AGL

Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.

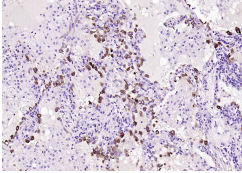
Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.

Background: FASN (Fatty Acid Synthase) is a central enzyme in de novo lipogenesis. FAS is a target for SREBP and is upregulated by LXR activation; it is also one of the accepted markers for insulin resistance, SREBP and LXR activation.

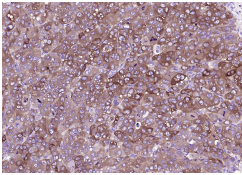
VALIDATION IMAGES



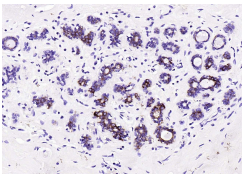
25 ug total protein per lane of various lysates (see on figure) probed with FASN monoclonal antibody, unconjugated (bsm-51155M) at 1:4000 dilution and 4°C overnight incubation. Followed by conjugated secondary antibody incubation at r.t. for 60 min.



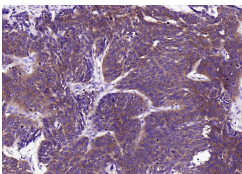
Paraformaldehyde-fixed, paraffin embedded (human lung carcinoma); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Incubation with (FASN) Monoclonal Antibody, Unconjugated (bsm-51155M) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Mouse)(sp-0024) instructions and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (human liver carcinoma); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Incubation with (FASN) Monoclonal Antibody, Unconjugated (bsm-51155M) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Mouse)(sp-0024) instructions and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (human breast); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Incubation with (FASN) Monoclonal Antibody, Unconjugated (bsm-51155M) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Mouse)(sp-0024) instructions and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (human breast carcinoma); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Incubation with (FASN) Monoclonal Antibody, Unconjugated (bsm-51155M) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Mouse)(sp-0024) instructions and DAB staining.

PRODUCT SPECIFIC PUBLICATIONS

[IF=3.654] Yingjun Zhou. et al. Chrysin improves diabetic nephropathy by regulating the AMPK-mediated lipid metabolism in HFD/STZ-induced DN mice. J FOOD BIOCHEM. 2022 Aug;;e14379 WB ; Mouse . 35976957

[IF=3.448] Yang X et al. miR - 760 exerts an antioncogenic effect in esophageal squamous cell carcinoma by negatively driving fat metabolism via targeting c - Myc. J Cell Biochem. 2019 Nov 10. WB ; Human . 31709636

[IF=3.231] Jing Fan. et al. Chitosan Oligosaccharide Inhibits the Synthesis of Milk Fat in Bovine Mammary Epithelial Cells through AMPK-Mediated Downstream Signaling Pathway. ANIMALS. 2022 Jan;12(13):1692 WB ; Bovine . 35804595