bs-18037R

[Primary Antibody]



OTC/Ornithine Carbamoyltransferase Rabbit pAb A N T | B

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DATASHEET -

Host: Rabbit Isotype: IgG

Clonality: Polyclonal

GeneID: 5009 **SWISS:** P00480

Target: OTC/Ornithine Carbamoyltransferase

Immunogen: KLH conjugated synthetic peptide derived from human

OTC/Ornithine Carbamoyltransferase: 31-130/354.

Purification: affinity purified by Protein A

Concentration: 1mg/ml

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: This nuclear gene encodes a mitochondrial matrix enzyme.

Missense, nonsense, and frameshift mutations in this enzyme lead to ornithine transcarbamylase deficiency, which causes

hyperammonemia. Since the gene for this enzyme maps close to that for Duchenne muscular dystrophy, it may play a role in that

disease also. [provided by RefSeq, Jul 2008]

Applications: WB (1:500-2000)

IHC-P (1:100-500) IHC-F (1:100-500) **IF** (1:100-500)

Flow-Cyt (1ug/Test)

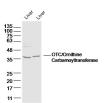
Reactivity: Human, Mouse, Rat

(predicted: Pig, Sheep, Cow, Dog, Horse)

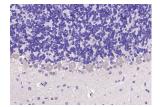
Predicted MW.: 36 kDa

Subcellular Cytoplasm

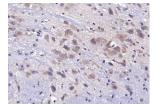
VALIDATION IMAGES -



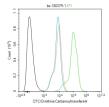
Sample: Liver (Mouse) Lysate at 40 ug Liver (Rat) Lysate at 40 ug Primary: Anti- OTC/Ornithine Carbamoyltransferase (bs-18037R) at 1/300 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 36 kD Observed band size: 36 kD



Paraformaldehyde-fixed, paraffin embedded (Rat cerebellum); 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; Antibody incubation with (OTC/Ornithine Carbamoyltransferase) Polyclonal Antibody, Unconjugated (bs-18037R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (mouse cerebellum); 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; Antibody incubation with (OTC/Ornithine Carbamoyltransferase) Polyclonal Antibody, Unconjugated (bs-18037R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Blank control (black line) :HepG2. Primary Antibody (green line): Rabbit Anti-Ornithine Carbamovltransferase antibody (bs-18037R) Dilution:1ug/Test; Secondary Antibody (white blue line): Goat anti-rabbit IgG-AF488 Dilution: 0.5ug/Test. Isotype control (orange line): Normal Rabbit IgG Protocol The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 90% ice-cold methanol for 20 min at -20°C, The cells were then incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at room temperature .Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.