**bsm-52458R** **[ Primary Antibody ]**

# ACLY Recombinant Rabbit mAb

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## DATASHEET

**Host:** Rabbit
**Isotype:** IgG
**Clonality:** Recombinant
**CloneNo.:** 3G8
**GeneID:** 47
**SWISS:** P53396
**Target:** ACLY

**Immunogen:** A synthesized peptide derived from human ATP citrate synthase: 1050-1101.

**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:** ATP citrate lyase is the primary enzyme responsible for the synthesis of cytosolic acetyl-CoA in many tissues. The enzyme is a tetramer (relative molecular weight approximately 440,000) of apparently identical subunits. It catalyzes the formation of acetyl-CoA and oxaloacetate from citrate and CoA with a concomitant hydrolysis of ATP to ADP and phosphate. The product, acetyl-CoA, serves several important biosynthetic pathways, including lipogenesis and cholesterogenesis. In nervous tissue, ATP citrate-lyase may be involved in the biosynthesis of acetylcholine. Two transcript variants encoding distinct isoforms have been identified for this gene. [provided by RefSeq]

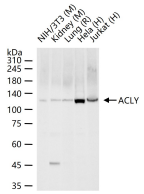
**Applications:** **WB** (1:500-2000)
**IHC-P** (1:50-200)
**IHC-F** (1:50-200)
**IF** (1:20-100)
**Flow-Cyt** (1:50-100)
**ICC/IF** (1:50-200)

**Reactivity:** Human, Mouse, Rat

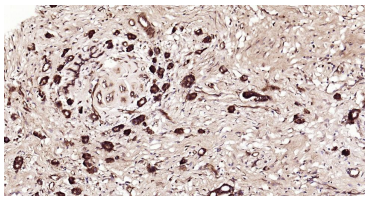
**Predicted MW.:** 122 kDa

**Subcellular Location:** Cytoplasm

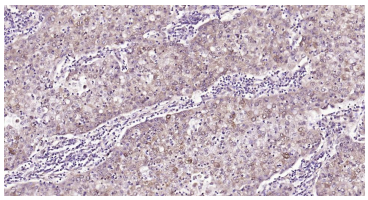
## VALIDATION IMAGES



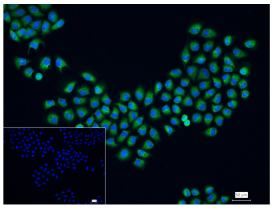

25 ug total protein per lane of various lysates (see on figure) probed with ACLY monoclonal antibody, unconjugated (bsm-52458R) at 1:2000 dilution and 4°C overnight incubation. Followed by conjugated secondary antibody incubation at r.t. for 60 min.



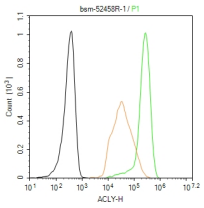
Paraformaldehyde-fixed, paraffin embedded Human Breast Cancer; Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15 min; Antibody incubation with ACLY Monoclonal Antibody, Unconjugated(bsm-52458R) at 1:200 overnight at 4°C, followed by conjugation to the SP Kit(Rabbit, SP-0023) and DAB (C-0010) staining.



Paraformaldehyde-fixed, paraffin embedded Human Gastric Cancer; Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15 min; Antibody incubation with ACLY Monoclonal Antibody, Unconjugated(bsm-52458R) at 1:200 overnight at 4°C, followed by conjugation to the SP Kit(Rabbit, SP-0023) and DAB (C-0010) staining.



4% Paraformaldehyde-fixed Hela (H) cell; Triton X-100 at r.t. for 20 min; Antibody incubation with (ACLY) monoclonal Antibody, unconjugated (bsm-52458R) 1:100, 90 min at 37°C; followed by conjugated Goat Anti-Rabbit IgG antibody (green, bs-40295G-FITC) at 37°C for 90 min, DAPI




The Hela (H) cells were fixed with 4% PFA (10 min at r.t.) and then permeabilized with 90% ice-cold methanol for 20 min at -20°C,the cells then were incubated in 5%BSA to block non-specific protein-protein interactions (30 min at r.t.).Primary Antibody (green):Rabbit Anti-ACLY

(blue, C02-04002) was used to stain the cell nuclei. PBS instead of the primary antibody was used as the blank control.

antibody (bsm-52458R,1:100); Isotype Control (orange): Rabbit IgG (bs-0295P). Blank control (black): PBS. Acquisition of 20,000 events was performed.