

**Manufacturer/Supplier:**

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**Product name:** Rabbit anti-ACLY Antibody

**Catalog:** DL93552A

**Synonyms:** ATP-citrate synthase; ATP-citrate (pro-S-)-lyase;  
ACL; Citrate cleavage enzyme

**Immunogen:** KLH-conjugated synthetic peptide encompassing a  
sequence within the center region of human ACLY.  
The exact sequence is proprietary.

**Form:** Liquid

**Concentration:** 1mg/mL

**Size:** 100 ul/50 ul

**Host:** Rabbit

**Reactivity:**  
Human, Mouse, Rat, Bovine, Pig

**Application:** ELISA, WB, IHC,  
IF/IC

**Clonality:** Polyclonal

**Dilution:** WB (1/500 - 1/1000),  
IHC (1/100 - 1/200), IF/IC (1/100 -  
1/500), IP (1/10 - 1/100)

**Entrez Gene:** 9212/20877/114592

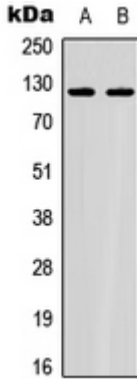
**SwissProt:** Q96GD4/O70126/O55099

**Purification:** The antibody was purified by immunogen affinity chromatography.

**Buffer:** Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.

**WB description:**

Western blot analysis of Aurora B expression in HepG2 (A), Raw264.7 (B), rat liver (C) whole cell lysates.

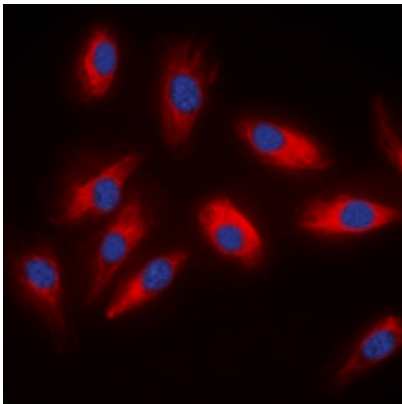


### IHC description:

Immunohistochemical analysis of Aurora B staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

### IF/ICC description:

Immunofluorescent analysis of Aurora B staining in HepG2 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antib



**Storage:**

Store at -20°C. Avoid repeated freeze / thaw cycles.