

BOSTER BIOLOGICAL TECHNOLOGY

Building C21, 3rd to 5th Floors, Optics Valley Biopharmaceutical Accelerator, East Lake High-Tech Development Zone, Wuhan.

Web: www.boster.com Phone: 027-67845390/1/2 Email: boster@boster.com

Basic Information	
Product Name	Anti-Gelsolin/GSN Antibody (Clone#GS-2C4)
Gene Name	GSN
Source	Mouse
Clonality	Monoclonal
lsotype	lgG1
Species Reactivity	human, rabbit
Tested Application	WB, IHC
Contents	200ug/ml antibody with PBS $ ightarrow$ 0.02% NaN3 , 1mg BSA and 50% glycerol.
Immunogen	Human plasma gelsolin.
Concentration	200ug/ml
Purification	Ascites
Observed MW	86 kDa
Dilution Ratios	Western blot (WB):1:500-2000Immunohistochemistry (IHC):1:50-400(Boiling the paraffin sections in 10mM citrate buffer,pH6.0,or PH8.0 EDTA repair liquid for 20mins is required for the staining of formalin/paraffin sections.) Optimal working dilutions must be determined by end user.

Storage

12 months from date of receipt, -20°C as supplied.

Background Information

Gelsolin, a protein of leukocytes, platelets, and other cells, severs actin filaments in the presence of submicromolar calcium, thereby solating cytoplasmic actin gels. A gelsolin variant with 23 more N-terminal amino acids is a plasma component probably involved in the clearance of actin, the most abundant human protein, from the circulation. Gelsolin is located in 9q34. Plasma and cytoplasmic gelsolins are encoded by a single gene and contain a duplicated actin-binding domain.

Selected Validation Data

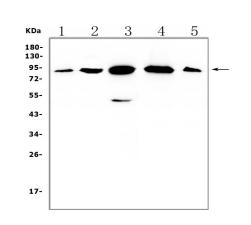
Product datasheet Anti-Gelsolin/GSN Antibody (Clone#GS-2C4) Catalog Number: BM1620



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Western blot analysis of Gelsolin using anti-Gelsolin antibody (BM1620). Electrophoresis was performed on a 5-20% SDS-PAGE gel at 70V (Stacking gel) / 90V (Resolving gel) for 2-3 hours. The sample well of each lane was loaded with 50ug of sample under reducing conditions.

Lane 1: human Hela whole cell lysate,

Lane 2: human A549 whole cell lysate,

Lane 3: monkey COS-7 whole cell lysate,

Lane 4: human Caco-2 whole cell lysate,

Lane 5: human HepG2 whole cell lysate.

After Electrophoresis, proteins were transferred to a Nitrocellulose membrane at 150mA for 50-90 minutes. Blocked the membrane with 5% Non-fat Milk/ TBS for 1.5 hour at RT. The membrane was incubated with mouse anti-Gelsolin antigen affinity purified monoclonal antibody (Catalog # BM1620) at 0.5 µg/mL overnight at 4°C, then washed with TBS-0.1%Tween 3 times with 5 minutes each and probed with a goat antimouse IgG-HRP secondary antibody at a dilution of 1:10000 for 1.5 hour at RT. The signal is developed using an Enhanced Chemiluminescent detection (ECL) kit (Catalog # EK1001) with Tanon 5200 system. A specific band was detected for Gelsolin at approximately 90KD. The expected band size for Gelsolin is at 86KD.