BOSTER BIOLOGICAL TECHNOLOGY Building C21, 3rd and 4th floors, Optics Valley Biomedical Accelerator, Wuhan East Lake High-tech Development Zone

ntibody and ELIS

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

| Basic Inform | ation | |
|---------------------|--|---|
| Product Name | Anti-c-Jun/JUN Antibody | |
| Gene Name | JUN | |
| Source | Rabbit | |
| Clonality | Polyclonal | |
| lsotype | IgG | |
| Species Reactivity | human | |
| Tested Application | WB, IHC, ICC/IF | |
| Contents | 500 ug/ml antibody with PBS, 0.02% NaN3, 1 mg/ml BSA and 50% glycerol. | |
| Immunogen | A synthetic peptide corresponding to a sequence in the middle region of human c- Jun/JUN, which shares 93.3% amino acid (aa) sequence identity with both mouse and rat c-Jun/JUN. | |
| Concentration | 500 ug/ml | |
| Purification | Immunogen affinity purified. | |
| Observed MW | 36-48 kDa | |
| Dilution Ratios | Western blot (WB): Immunohistochemistry (IHC): Immunocytochemistry/Immunofluorescence (ICC/IF): (Boiling the paraffin sections in 10mM citrate buffer,pH6.0,o for 20 mins is required for the staining of formalin/paraffin s dilutions must be determined by end user. | 1:500-2000 1:50-400 1:50-400 r PH8.0 EDTA repair liquid ections.) Optimal working |

Storage

12 months from date of receipt, -20°C as supplied. 6 months 2 to 8°C after reconstitution. Avoid repeated freezing and thawing.

Background Information

c-Jun is a protein that in humans is encoded by the JUN gene. This gene is the putative transforming gene of avian sarcoma virus 17. It encodes a protein which is highly similar to the viral protein, and which interacts directly with specific target DNA sequences to regulate gene expression. This gene is intronless and is mapped to 1p32-p31, a chromosomal region involved in both translocations and deletions in human malignancies.

antibody and ELISA experts BOSTER BIOLOGICAL TECHNOLOGY

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Selected Validation Data



Figure 1. Western blot analysis of anti- JUN Antibody (A02038-2). The sample well of each lane was loaded with 50ug of sample under reducing conditions. Lane 1: U2OS whole cell lysates, Lane 2: PC-3 whole cell lysates. Use rabbit anti- JUN 1:1000, probed with a goat anti-rabbit IgG-HRP secondary antibody. The signal is developed using an Enhanced Chemiluminescent detection (ECL) kit (Catalog # EK1002). A specific band was detected for JUN at approximately 43KD. The expected band size for JUN is at 36KD.



Figure 2. IHC analysis using anti- JUN Antibody (A02038-2). detected in paraffin-embedded section of human lung cancer tissue. Biotinylated goat anti-rabbit IgG was used as secondary antibody. The tissue section was developed using Strepavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB as the chromogen.



Figure 8. ICC analysis using anti- JUN Antibody (A02038-2). was detected in immersion fixed A431 cell line. Cells were stained using the Dylight488-conjugated Anti-rabbit IgG Secondary Antibody (green)(Catalog # BA1127) and counterstained with DAPI (blue).