

Technical Data Sheet

Purified Mouse Anti-Jun

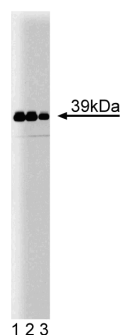
Product Information

| | |
|-------------------------|--|
| Material Number: | 610326 |
| Size: | 50 µg |
| Concentration: | 250 µg/ml |
| Clone: | 3/Jun |
| Immunogen: | Mouse c-Jun aa. 26-175 |
| Isotype: | Mouse IgG2a |
| Reactivity: | QC Testing: Human Tested in Development: Chicken, Dog, Mouse, Rat |
| Target MW: | 39 kDa |
| Storage Buffer: | Aqueous buffered solution containing BSA, glycerol, and ≤0.09% sodium azide. |

Description

The activator protein transcription factor (AP-1) was identified as a protein that recognizes specific sequences in the *cis*-control regions of the SV40 virus and the human metallothionein IIA gene. AP-1 is composed of protein products of two different gene families: *jun* and *fos*. The AP-1 transcription factor is either a homodimer of Jun proteins or a heterodimer of Jun and Fos proteins. The transcriptional activity of Jun is enhanced by phosphorylation in its activation domain at Ser63 and Ser73. Phosphorylation at both sites is necessary for stimulating the activating function of Jun. Jun is phosphorylated by JNK protein kinases that are activated by the same signals that potentiate Jun activity.

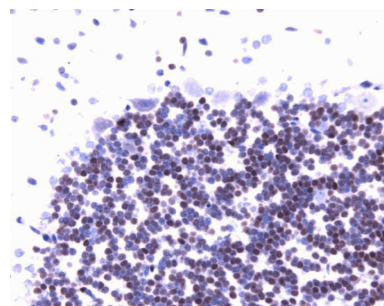
This antibody is routinely tested by western blot analysis. Other applications were tested at BD Biosciences Pharmingen during antibody development only or reported in the literature.



Western blot analysis of Jun on human endothelial cell lysate. Lane 1: 1:1000, lane 2: 1:2000, lane 3: 1:4000 dilution of anti-Jun antibody.



Immunohistochemical staining of Rabbit Muscle tissue section. SDS-treated formalin-fixed paraffin-embedded section.



Immunohistochemical staining of rat brain. Formalin-fixed paraffin-embedded section with citrate buffer pretreatment. 40X.

Preparation and Storage

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography.

Store undiluted at -20° C.

Application Notes

Application

| | |
|----------------------|---------------------------|
| Western blot | Routinely Tested |
| Immunofluorescence | Tested During Development |
| Immunohistochemistry | Tested During Development |
| Immunoprecipitation | Tested During Development |

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Suggested Companion Products

| Catalog Number | Name | Size | Clone |
|----------------|-------------------------------|--------|------------|
| 611450 | Human Endothelial Cell Lysate | 500 µg | (none) |
| 554002 | HRP Goat Anti-Mouse Igs | 1.0 ml | (none) |
| 554001 | FITC Goat Anti-Mouse Igs | 0.5 mg | Polyclonal |

Product Notices

1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
2. Please refer to www.bdbiosciences.com/pharming/en/protocols for technical protocols.
3. Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before discarding to avoid accumulation of potentially explosive deposits in plumbing.
4. Source of all serum proteins is from USDA inspected abattoirs located in the United States.

References

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Lamph WW, Wamsley P, Sassone-Corsi P, Verma IM. Induction of proto-oncogene JUN/AP-1 by serum and TPA. *Nature.* 1988; 334(6183):629-631.(Biology)