Technical Data Sheet

Purified Mouse Anti-JNK/SAPK (pT183/pY185)

Product Information

Material Number:	612540
Size:	50 µg
Concentration:	250 µg/ml
Clone:	41/JNK/SAPK (pT183/pY185)
Immunogen:	Phosphorylated Human JNK/SAPK (pT183/pY185) Peptide
Isotype:	Mouse IgG1
Reactivity:	QC Testing: Human
	Tested in Development: Mouse, Rat
Target MW:	43 & 56 kDa
Storage Buffer:	Aqueous buffered solution containing BSA, glycerol, and ≤0.09% sodium
	azide.

Description

The Ras signaling pathway links the signals from growth factor receptors with the activation of the MAPK kinase cascade of phosphorylation leading to cell growth and differentiation. External stimuli, like endotoxins, UV irradiation, heat, and hyperosmolarity, induce an array of cellular responses that culminate with gene expression, ultimately dictating an adaptation to the new environment. Small GTPases of the Rho family, including cdc42, Rac1, and Rho, transmit the stress signals that initiate the signal cascade. JNK is a c-Jun kinase that was also identified as SAPK1 and MAPKp49. JNK/SAPK, along with p38 and RK5/BMK1, comprise three classes of stress-activated MAPK groups. Complete activation of JNK/SAPK requires the phosphorylation of both Thr183 and Tyr185, which are located in a Thr-X-Tyr motif. The activation of these residues is believed to be carried out by MKK4 and MKK7. Active JNK/SAPK phosphorylates other kinases and multiple transcription factors that induce expression of genes, such as proinflammatory cytokines.



Western blot analysis for JNK/SAPK (pT183/pY185). HeLa cells (Human cervical epitheloid carcinoma; ATCC CCL-2) were either left untreated (lane 1) or treated with 25 μ g/mL anisomycin, an antibiotic and protein synthesis inhibitor, for 15 min at 37°C (lane 2). The top panel was probed with a mouse anti-JNK/SAPK1 antibody (Cat. No. 610627) and the bottom panel was probed with the mouse anti-JNK/SAPK1 (pT183/pY185) antibody at a 1:250 dilution with bands observable at ~ 43 kDa & ~ 56 kDa.

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			
Flow cytometry	Not Recommended			

Recommended Assay Procedure:

Western blot: Please refer to http://www.bdbiosciences.com/pharmingen/protocols/Western_Blotting.shtml

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Suggested Companion Products

Catalog Number	Name	Size	Clone
611449	HeLa Cell Lysate	500 µg	(none)
611692	HeLa + Anisomycin Cell Lysate	500 μg	(none)
554002	HRP Goat Anti-Mouse Ig	1.0 ml	(none)
610627	Purified Mouse Anti-JNK/SAPK1	50 µg	37/pan-JNK/SAPK1

Product Notices

- 1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
- 2. Please refer to www.bdbiosciences.com/pharmingen/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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