

**Antibody Datasheet**

**Product description:-** Anti-TAB1 phospho Thr 431

**Antigen:-** CSTLDEAT\*PTLTNQ [\* is phospho Threonine]

**Sheep Number:-** S737A

**Bleed Number:-** Second Bleed

**Concentration:-** 0.34 mg/ml

**Formulation:-** Phosphate Buffered Saline

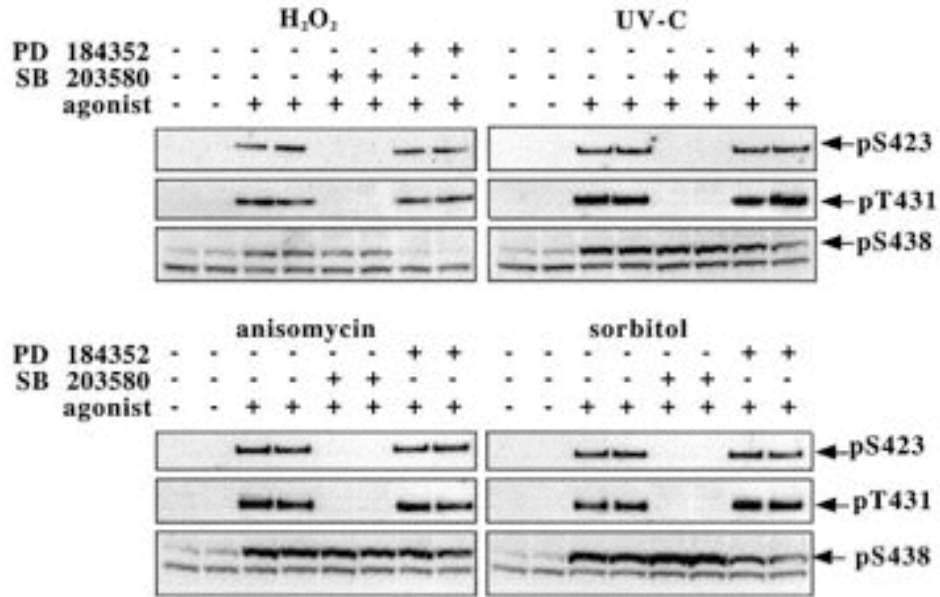
**Storage temperature:-** -20 °C

**Purification Method:-** Affinity purified against phospho peptide

**Working Concentration:-** 0.5-1  $\mu$ g/ml for western blotting  
Include the unphosphorylated form of the peptide immunogen at 10  $\mu$ g/ml to the diluted antibody

**Publication Reference:-**

“Feedback control of the protein kinase TAK1 by SAPK2a/p38alpha”  
Cheung PCF, Campbell DG, Nebreda AR and Cohen P.  
EMBO Journal. 2003, vol. 22 no. 21. 5793 - 5805.



KB cells were pre-treated for 1 hour with or without 10  $\mu$ M SB 203580 or 2  $\mu$ M PD184352 then simulated with H<sub>2</sub>O<sub>2</sub> (2 mM for 15 mins), UV-C (30 mins), anisomycin (10  $\mu$ g/ml for 30 mins) or sorbitol (0.5 M for 30 mins) and then lysed. TAB1 was immunoprecipitated using anti-TAB1 S823A, 2  $\mu$ g of anti-TAB1 was incubated for 1 hour at 4 °C with 250  $\mu$ g of protein lysate. The immunocomplex was denatured in SDS, separated by SDS-PAGE, transferred to nitrocellulose membrane and blotted with the appropriate phospho-specific antibody.

Binding of the primary antibody was detected using rabbit peroxidase conjugated anti-sheep IgG antibody (1 in 10, 000 dilution, Pierce) followed by enhanced chemiluminescence (ECL, Amersham).