

# Product Datasheet

## HSP27 [p Ser86] Antibody NSB536

Unit Size: 0.1 ml

Aliquot and store at -20C or -80C. Avoid freeze-thaw cycles.

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**NSB536****HSP27 [p Ser86] Antibody**

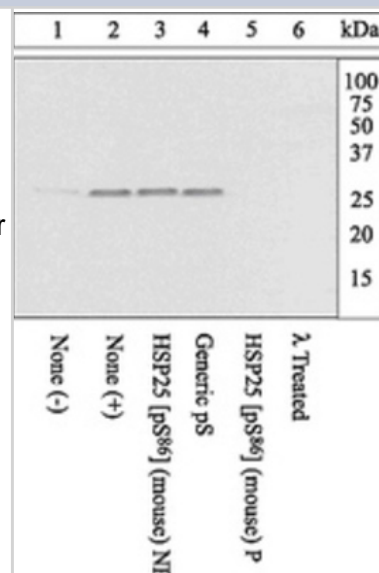
Product Information	
<b>Unit Size</b>	0.1 ml
<b>Concentration</b>	Please see the vial label for concentration. If unlisted please contact technical services.
<b>Storage</b>	Aliquot and store at -20C or -80C. Avoid freeze-thaw cycles.
<b>Clonality</b>	Polyclonal
<b>Preservative</b>	0.05% Sodium Azide
<b>Purity</b>	Immunogen affinity purified
<b>Buffer</b>	Dulbecco's PBS (pH 7.3), 1.0 mg/ml and 50% Glycerol

Product Description	
<b>Host</b>	Rabbit
<b>Gene ID</b>	3315
<b>Gene Symbol</b>	HSPB1
<b>Species</b>	Mouse
<b>Reactivity Notes</b>	Mouse HSP25. Endogenous human HSP27 phosphorylated at serine 82 (HeLa cells treated with TNF-a) was weakly detected by this antibody.
<b>Specificity/Sensitivity</b>	HSP25 Phosphospecific [Ser86]
<b>Immunogen</b>	The antiserum was produced against a chemically synthesized phosphopeptide derived from a region of mouse HSP25 that contains serine 86.

Product Application Details	
<b>Applications</b>	Western Blot
<b>Recommended Dilutions</b>	Western Blot
<b>Application Notes</b>	The antibody has been used for Western blotting 1:1000. The optimal antibody concentration should be determined empirically for each specific application.

**Images**

Peptide Competition Lysates prepared from NIH3T3 cells left unstimulated (1) or treated with anisomycin (2-6) were resolved by SDS-PAGE on a 10% polyacrylamide gel and transferred to PVDF. Membranes were left untreated (1-5) or treated with lambda (l) phosphatase (6), blocked with a 5% BSA-TBST buffer for one hour at room temperature, and incubated with HSP25 [pS86] (mouse) antibody for one hour at room temperature in 3% BSA-TBST buffer, following prior incubation with: no peptide (1, 2, 6), the non-phosphopeptide corresponding to the immunogen (3), a generic phosphoserine-containing peptide (4), or, the phosphopeptide immunogen (5). After washing, membranes were incubated with goat F(ab)2 anti-rabbit IgG HRP conjugate in 3% BSA-TBST buffer, and bands were detected using the Pierce SuperSignal(TM) method. The data show that only the peptide corresponding to HSP25 [pS86] (mouse) blocks the antibody signal, thereby demonstrating the specificity of the antibody. The signal was completely removed by l phosphatase treatment demonstrating that the antibody interacts specifically with the phosphorylated protein.





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### **Limitations**

This product is for research use only and is not approved for use in humans or in clinical diagnosis. Primary Antibodies are guaranteed for 1 year from date of receipt.

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