Human Small Ribonucleoprotein Associated Protein B B' (SmB/B')

Origin:	Recombinant	Cat. No.:	41530
Tag:	N-terminal 6xHis	Size:	0.1 mg
Source:	Spodoptera frugiperda Sf9	Purity:	>90%
Other Names:	SmB/B', snRNPB/B'	Species:	Human

Description

Expressed in insect Sf9 cells with total 265 AA. Mw: 27.6 KDa (calculated). N-terminal 6xHis-tag and TEV cleavage site, 25 extra AA (highlighted). **Recombinant antigen for research use or manufacturing only.**

Introduction to the Molecule

Small nuclear ribonucleoprotein complexes (abbreviated as U-snRNP) are essential for splicing of precursor mRNA molecules and snRNPB is one of several nuclear proteins which are found in common among U1, U2, U4/U6, and U5 small ribonucleoprotein particles (snRNPs). Autoantibodies from patients with systemic lupus erythematosus (SLE) recognize epitopes on snRNPB/B'.

Immunological Function

As an autoantigen, SmB/B' binds with IgG-type human auto-antibodies.

Amino Acid Sequence

MSYYHHHHHHDYDIPTTENLYFQGATVGKSSKMLQHIDYRMRCILQDGRIFIGTFKAFDKHMNLI LCDCDEFRKIKPKNSKQAEREEKRVLGLVLLRGENLVSMTVEGPPPKDTGIARVPLAGAAGGPGI GRAAGRGIPAGVPMPQAPAGLAGPVRGVGGPSQQVMTPQGRGTVAAAAAAATASIAGAPTQYPPG RGGPPPPMGRGAPPPGMMGPPPGMRPPMGIPPGRGTPMGMPPPGMRPPPPGMRGPPPPGMR PPRP.

Applications

Standard ELISA test, line/dot assay and microarray assay with positive/negative sera panels.

Formulation

Lyophilized at 1 mg/mL in storage buffer (20mM Tris, 300-500mM NaCl, 10% Glycerol, PH8.0).

Reconstitution

Add deionized water to prepare a working stock solution of approximately 1 mg/mL and let the lyophilized pellet dissolve completely.

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Storage

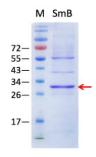
IMD

Store lyophilized protein at -20°C. Aliquot reconstituted protein and store at -80°C. Avoid repeated freezing /thawing cycles.

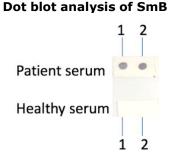
Quality Control Test

BCA to determine quantity of the protein. SDS PAGE to determine purity of the protein. Immunodot analysis to determine functionality of protein.

SDS-PAGE Gel



Dot blot assay



Analysis of serum from healthy subjects and patients. Recombinant autoantigens were utilized in this dot-blot assay for validation

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