20S Immunoproteasome, mouse Spleen

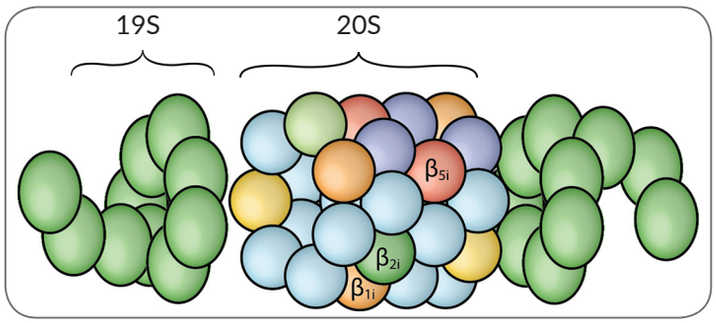
Cat. No. SBB-PP0083
Lot. No. 190700083

South Bay Bío

20S Immunoproteasome

The immunoproteasome is structurally similar to constitutive 26S proteasome. The 20S core of immunoproteasome contains two outer rings composed of alpha subunits, and two internal 7-subunit containing rings each possessing 3 specific subunits responsible for proteasome catalytic activity. In immunoproteasome these subunits (ß1, ß2, ß5) are replaced by three inducible subunits: PSMB9, PSMB10, and PSMB8, (ß1i, ß2i, ß5i). These stress-induced subunits allow for the production of MHC-1 associating peptides, which are displayed as antigens on the cell surface. These displayed peptides can then be recognized by immune surveillance CD8 T-Cells. 20S

Immunoproteasome is recognized as a strong drug target for autoimmune disease and cancer. This immunoproteasome is purified from mouse spleen and is supplied at >95% purity. The Immunoproteasome is commonly associated with the 19S, PA28 α/ß, or the PA28γ regulatory complexes. If choosing to omit PA28 during use, 20S must be chemically activated by addition of 0.035%SDS in final assay buffers. Optimal eperimental concentrations are between 2-5 nM. The relatively promient band at ~90 kDa is associated to Hsp90 and commonly associated with the proteasome.



Product Information

Quantity: 25 µg Molecular Weight: >700 kDa

Concentration: 1.6 µM, 1.1 mg/mL

Purity: >92% by SDS-PAGE

Storage Buffer: 50 mM HEPES pH 7.5, 100 mM NaCl, 1 mM TCEP.

Storage: Store at -80˚C. Avoid multiple freeze thaw-cycles

Quality Control and Performance

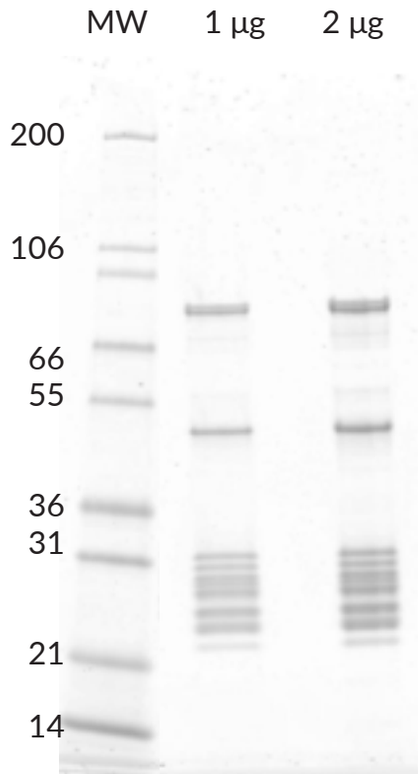


Figure 1. 20S Immunoproteasome, SDS-PAGE. From left to right, increasing amounts of 20S Immunoproteasome were loaded onto a 4-20% SDS-PAGE gel, and stained with Coomassie brilliant blue.

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www.southbaybio.com

Contact:
info@southbaybio.com

5941 Optical Ct, Suite 229
San Jose, CA 95138 USA

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Quality Control and Performance Data

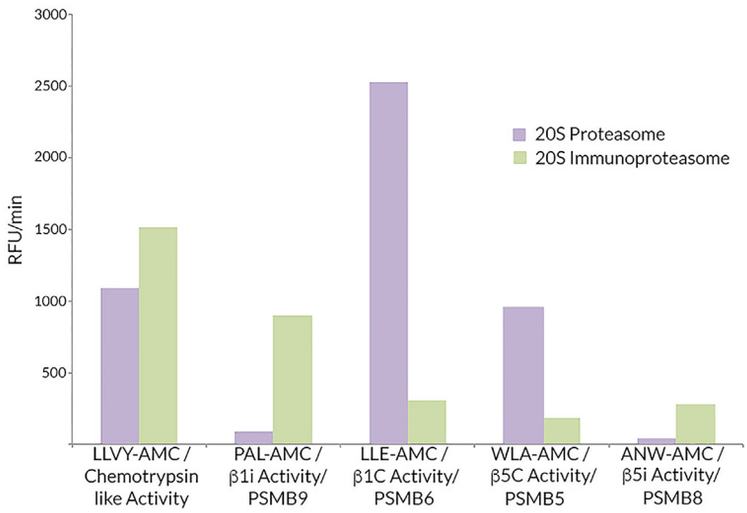

Figure 2. 20S Immunoproteasome vs. 20S Constitutive Proteasome Activity. 20S Immunoproteasome is most active against LLVY-AMC (SBB-PS0010), PAL-AMC (SBB-PS0007), and ANW-AMC (SBB-PS0009) substrates, representing physiologically relevant chemotrypsin-like, ß1i, and ß5i immunoproteasome activity respectively.

References

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3)Cascio P, Hilton C, Kisselev AF, Rock KL, Goldberg AL (May 2001). "26S proteasomes and immunoproteasomes produce mainly N-extended versions of an antigenic peptide". The EMBO Journal. 20 (10): 2357–66. doi:10.1093/emboj/20.10.2357. PMC 125470free to read. PMID 11350924.

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