

# 20S Immunoproteasome, rat Spleen

Cat. No. SBB-PP0046  
Lot. No. 190700046



# South Bay Bio

## 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 ( $\beta$ 1,  $\beta$ 2,  $\beta$ 5) are replaced by three inducible subunits: PSMB9, PSMB10, and PSMB8, ( $\beta$ 1i,  $\beta$ 2i,  $\beta$ 5i). 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 rat spleen and is supplied at >95% purity. The Immunoproteasome is commonly associated with the 19S, PA28  $\alpha/\beta$ , or the PA28 $\gamma$  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 experimental concentrations are between 2-5 nM.

## Product Information

Quantity: 25  $\mu$ g      Molecular Weight: >700 kDa

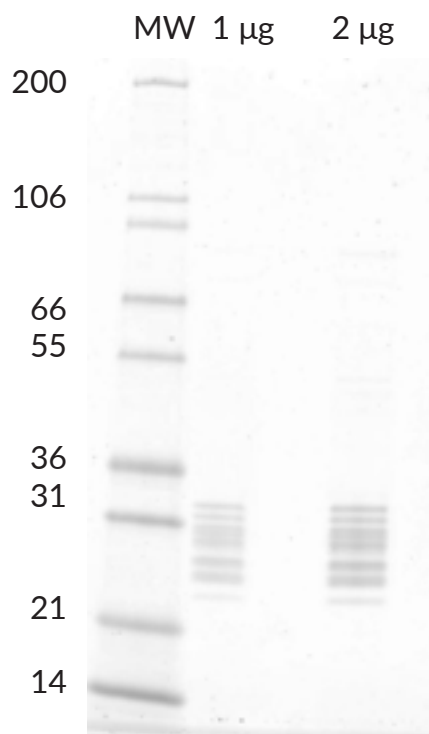
Concentration: 1.6  $\mu$ M, 1.1 mg/mL

Purity: >95% by SDS-PAGE

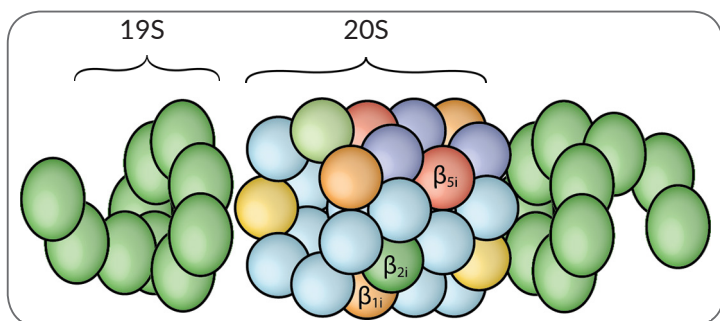
Storage Buffer: 50 mM HEPES pH 7.5, 100 mM NaCl, 1 mM TCEP.

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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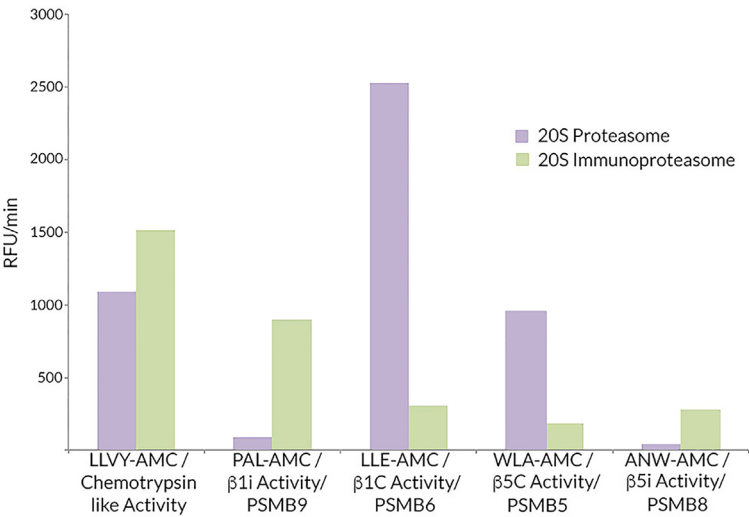
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## Quality Control and Performance Data



**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 chemo-trypsin-like,  $\beta$ 1i, and  $\beta$ 5i immunoproteasome activity respectively.

## References

1) Wang J, Maldonado MA (Aug 2006). "The ubiquitin-proteasome system and its role in inflammatory and autoimmune diseases". Cellular & Molecular Immunology. 3 (4): 255-61. PMID 16978533.

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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 125470 free to read. PMID 11350924.

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