

## DESCRIPTION

**Source** *Spodoptera frugiperda*, Sf21 (baculovirus)-derived  
 Leu2-Gln710, with an N-terminal Met and 6-His tag  
 Accession # Q9QUR6

**N-terminal Sequence Analysis** Met

**Predicted Molecular Mass** 82 kDa

## SPECIFICATIONS

**SDS-PAGE** 68-73 kDa, reducing conditions

**Activity** Measured by its ability to convert the substrate benzyloxycarbonyl-Gly-Pro-7-amido-4-methylcoumarin (Z-GP-AMC) to Z-Gly-Pro and 7-amino-4-methylcoumarin (AMC).  
 The specific activity is >4,500 pmol/min/μg, as measured under the described conditions. See Activity Assay Protocol on [www.RnDSystems.com](http://www.RnDSystems.com)

**Endotoxin Level** <1.0 EU per 1 μg of the protein by the LAL method.

**Purity** >90%, by SDS-PAGE under reducing conditions and visualized by Colloidal Coomassie® Blue stain at 5 μg per lane.

**Formulation** Supplied as a 0.2 μm filtered solution in MES, NaCl and Glycerol. See Certificate of Analysis for details.

## Activity Assay Protocol

- Materials**
- Assay Buffer: 25 mM Tris, 250 mM NaCl, 2.5 mM DTT, pH 7.5
  - Recombinant Mouse Prolyl Oligopeptidase/PREP (rmPREP) (Catalog # 6339-SE)
  - Substrate: Z-GP-AMC (Bachem, Catalog # I-1145), 10 mM stock in DMSO
  - F16 Black Maxisorp Plate (Nunc, Catalog # 475515)
  - Fluorescent Plate Reader (Model: SpectraMax Gemini EM by Molecular Devices) or equivalent

- Assay**
1. Dilute rmPREP to 0.1 ng/μL in Assay Buffer.
  2. Dilute Substrate to 100 μM in Assay Buffer.
  3. Load into a plate 50 μL of 0.1 ng/μL rmPREP, and start the reaction by adding 50 μL of 100 μM Substrate.
  4. For Substrate Blanks, load 50 μL of Assay Buffer and 50 μL of 100 μM Substrate.
  5. Read plate at excitation and emission wavelengths of 380 nm and 460 nm, respectively, in kinetic mode for 5 minutes.
  6. Calculate specific activity:

$$\text{Specific Activity (pmol/min/}\mu\text{g)} = \frac{\text{Adjusted } V_{\text{max}}^* \text{ (RFU/min)} \times \text{Conversion Factor}^{**} \text{ (pmol/RFU)}}{\text{amount of enzyme (}\mu\text{g)}}$$

\*Adjusted for Substrate Blank

\*\*Derived using calibration standard 7-Amino, 4-Methyl Coumarin (Sigma, Catalog # A-9891).

- Final Assay Conditions**
- Per Well:
- rmPREP: 0.005 μg
  - Substrate: 50 μM

## PREPARATION AND STORAGE

**Shipping** The product is shipped with polar packs. Upon receipt, store it immediately at the temperature recommended below.

- Stability & Storage** **Use a manual defrost freezer and avoid repeated freeze-thaw cycles.**
- 6 months from date of receipt, -20 to -70 °C as supplied.
  - 3 months, -20 to -70 °C under sterile conditions after opening.

\* Coomassie is a registered trademark of Imperial Chemical Industries Ltd.

## BACKGROUND

Prolyl Oligopeptidase, also known as prolyl endopeptidase and post-proline cleaving enzyme, is a serine peptidase displaying specificity for the cleavage of Pro-Xaa bonds of oligopeptide substrates (1, 2). The peptidase is known to hydrolyze a variety of biologically active peptides such as bradykinin, substance P, neurotensin, and vasopressin (3). Because of its action on neuropeptides, Prolyl Oligopeptidase is considered to be involved in processes such as learning, memory, and depression (4).

## References:

1. Szeltner, Z and L. Polgár (2008) Current Protein Pept. Sci. **9**:96.
2. Yoshimoto, T. *et al.* (1977) Biochemistry. **16**:2942.
3. Wilk, S. (1983) Life Sci. **33**:2149.
4. Männistö, P.T. *et al.* (2007) Drug News Perspect. **20**:293.