

# SuperNuclease

catalog Number : BD-PD266651



## General Information

### Gene Name Synonym:

Nuclease, *Serratia marcescens*' extracellular endonuclease

### Protein Construction:

SuperNuclease is a recombinant *Serratia marcescens*' extracellular endonuclease.

**Source:** *Serratia marcescens*

**Expression Host:** *E.coli*

## QC Testing

**Purity:** > 95 % as determined by SDS-PAGE.  
> 99 % as determined by SEC-HPLC Analysis.

### Endotoxin:

< 0.01 EU/1000 units protein as determined by the LAL method.

### Activity:

Measured by its ability to cleave Salmon sperm DNA substrate .  
One unit is defined as the amount of enzyme that causes a  $\Delta A_{260}$  of 1.0 in 30 minutes at 37° C.  
The specific activity is  $\geq 1.1 \times 10^6$  units/mg.

### Stability:

Samples are stable for up to twelve months from date of receipt at -70 °C

### Predicted N terminal:

Three isoforms with different N terminal may be found from the compound—Sm1 (22D-266N), Sm2 (23T-266N) and Sm3 (25E-266N), the activity analysis shows that they were functionally equivalent

### Molecular Mass:

The SuperNuclease comprises 266 amino acids and has a calculated molecular mass of Sm1 (26708.2 Da), Sm2 (26591.8 Da) and Sm3 (26376.4 Da). The apparent molecular mass of SuperNuclease is approximately 26.5 KDa.

## Usage Guide

### Storage:

Store it under sterile conditions at -20°C to -80°C upon receiving. Recommend to aliquot the protein into smaller quantities for optimal storage.

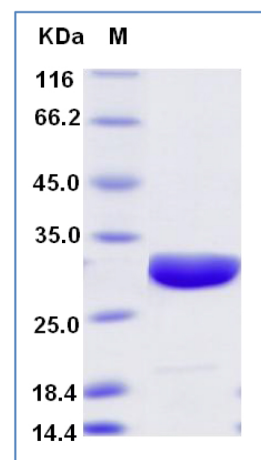
**Avoid repeated freeze-thaw cycles.**

## Protocol

### Large scale cell lysis treatment:

- 1). Lysis buffer preparation:  
The lysis buffer should be suitable for the protein as well as the downstream purification processes, and so on;
- 2). Resuspend cell plates in lysis buffer:  
The ratio of lysis buffer (mL) against the gram of cell can be (10-20):1;
- 3). Add SuperNuclease (250 units to 1 g cells):  
It is well recommended to optimize the amount of SuperNuclease;
- 4). Lysate cell by mechanical or chemical methods on ice or at room temperature:  
Methods: Ultrasonic disruption, High Pressure Homogenizer, tissue homogenizer, and so on;
- 5). Obtain clear cell lysate supernatant by centrifugation at ~12,000 rpm for 0.5 hour.

### SDS-PAGE:



### SEC-HPLC:

