

# Cas9 Nuclease protein NLS (3 µg/µl)

# Cas9 Nuclease protein NLS (15 µg/µl)

**保存:**

−20°C

**製品説明:**

本品は、*Streptococcus pyogenes* 由来の Cas9 Nuclease を、大腸菌で発現・精製した組換えタンパク質です。本品は核移行シグナル(NLS)を N 末端と C 末端に有しており、ガイド RNA と組み合わせることによりゲノム編集に利用することができます。

**製品内容:**
**Code No. 319-08641**

構成品	容量
Cas9 Nuclease protein NLS (3 µg/µL)	75 µg

**Code No. 316-08651**

構成品	容量
Cas9 Nuclease protein NLS (15 µg/µL)	300 µg

**形状:**

10 mM Tris-HCl (pH 7.5), 300 mM NaCl, 0.1 mM EDTA, 1 mM DTT, 50 % Glycerol

**起源:**

遺伝子組換え大腸菌

**分子量 × 10<sup>-3</sup>:**

162.75

**純度:**

- ・ 本酵素 10 µg と基質 DNA を 37°C で 1 時間反応させても、DNA のアガロースゲル電気泳動パターンに変化は認められない。
- ・ 本酵素 10 µg と基質 RNA を 37°C で 1 時間反応させても、RNA のアガロースゲル電気泳動パターンに変化は認められない。
- ・ エンドキシン < 1 EU/µg

**使用例:**

<in vitro>での切断チェック>

Cas9 Nuclease protein NLS 150 - 300 ng  
(約 1-2 pmol)

tracrRNA (2 µM) 1 µL  
crRNA (2 µM) 1 µL  
↓  
室温, 5 min.  
↓  
基質 DNA 100 ng  
10x H Buffer\*1) 2 µL  
-----  
d.d.H<sub>2</sub>O up to 20 µL  
↓  
37°C, 60 min.  
↓  
Loading Buffer (SDS 含有)\*2)  
↓  
65°C, 5 min.  
↓  
アガロースゲル電気泳動

\*1) 組成: 500 mM Tris-HCl (pH 7.9), 1 M NaCl, 100 mM MgCl<sub>2</sub>, 10 mM DTT

\*2) 正確な泳動結果を得るため、SDS を含む Loading Buffer (Code No.313-90111 など) の使用をお勧めします。

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