## Nuclease ELISA Kit

| Catalog Number | LDG00021E |
| :--- | :--- |
| Package | 96 T $(8 \times 12$ strips $) /$ Customized package |

For full product information, images and publications, please visit our website.

## Overview

## Description

Nuclease is a genetically engineered endonuclease from Serratia marcescens, it is produced and purified from E . coli. The enzyme is a non-specific nuclease, which effectively degrades single-stranded, double-stranded, linear, and circular DNA and RNA over a wide temperature and pH range.

Components

| Nuclease ELISA plate | 96 wells ( $12 \times 8$-well strips) |
| :--- | :---: |
| Standard | 1 vial (Lyophilized form) |
| Standard reconstitution buffer | 1 vial (1.5 mL) |
| HRP-antibody conjugate | 1 vial ( $70 \mathrm{\mu L}$ ) |
| HRP-antibody conjugated diluent buffer | 1 vial (12 mL) |
| $20 \times$ wash buffer | 1 vial ( 15 mL$)$ |
| TMB | 1 vial ( 12 mL$)$ |
| Stop solution | 1 vial ( 6 mL$)$ |
| Microplate sealing film | 1 sheet |

Specifications

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## Reactivity

Serratia marcescens

## Sensitivity

Limit of detection (LoD): $0.062 \mathrm{ng} / \mathrm{mL}$.
Limit of quantification (LoQ): $0.163 \mathrm{ng} / \mathrm{mL}$.

## Application

Sandwich ELISA analysis

## Specificity

Nuclease

## Instruction

## Shipping

The product is shipped with polar packs. Upon receipt, store it immediately at $2-8^{\circ} \mathrm{C}$ for long term storage.

## Stability \& Storage

This product is stable after storage at:

- $2-8^{\circ} \mathrm{C}$ for unopened product.

Please refer to product manual for storage constructions.

## Image

\section*{| Standard | Nuclease (ng/mL) | OD |  |
| :---: | :---: | :---: | :---: |
| 1 | 5 | 2.47 |  |
| 2 | 2.5 | 1.316 | 2.563 |
| 3 | 1.25 | 0.673 | 0.26 |
| 4 | 0.625 | 0.37 | 0.39 |
| 5 | 0.3125 | 0.229 | 0.234 |
| 6 | 0.15625 | 0.154 | 0.161 |
| 7 | 0.078125 | 0.128 | 0.117 |
| Blank | Blank | 0.093 | 0.083 | <br> }

Typical data
The following data are for demonstration only

|  | 1 | 2 | 3 | 4 |
| :---: | :---: | :---: | :---: | :---: |
| $A=N=$ | Standard 1 ( $5 \mathrm{ng} / \mathrm{mL}$ ) | Standard 1 ( $5 \mathrm{ng} / \mathrm{mL}$ ) | Sample 1 | Sample 5 |
| 8 | Standard 2 ( $2.5 \mathrm{ng} / \mathrm{mL}$ ) | Standard 2 ( $2.5 \mathrm{ng} / \mathrm{mL}$ ) | Sample 1 | Sample 5 |
| c | Standard 3 $(1.25 \mathrm{ng} / \mathrm{mL})$ | $\begin{gathered} \text { Standard } 3 \\ (1.25 \mathrm{ng} / \mathrm{mL}) \\ \hline \end{gathered}$ | Sample 2 | Sample 6 |
| ${ }^{\circ}$ | Standard 4 $(0.625 \mathrm{ng} / \mathrm{mL})$ | Standard 4 ( $0.625 \mathrm{ng} / \mathrm{mL}$ ) | Sample 2 | Sample 6 |
| E | $\begin{gathered} \text { Standard } 5 \\ (0.3125 \mathrm{ng} / \mathrm{mL}) \end{gathered}$ | $\begin{gathered} \text { Standard 5 } \\ (0.3125 \mathrm{ng} / \mathrm{mL}) \\ \hline \end{gathered}$ | Sample 3 | Sample 7 |
| F | $\begin{array}{\|c\|} \hline \text { Standard } 6 \\ (0.15625 \mathrm{ng} / \mathrm{mL}) \end{array}$ | $\begin{array}{\|c} \hline \text { Standard } 6 \\ (0.15625 \mathrm{ng} / \mathrm{mL}) \\ \hline \end{array}$ | Sample 3 | Sample 7 |
| ${ }^{6}$ ENE | Standard 7 $(0.078125 \mathrm{ng} / \mathrm{mL})$ | $\begin{gathered} \text { Standard 7 } \\ 0.078125 \mathrm{ng} / \mathrm{mLL} 2 \end{gathered}$ | $\text { Sample } 4$ | Sample 8 |
| H | Blank | ${ }^{\text {Blank }}$ | Sample 4 | Sample 8 |

An example of orientation of standards, blanks and samples in the stripwells microplate

| $\begin{gathered} \text { Reagents } \\ \text { (Store at } 2.8^{\circ} \mathrm{C} \text { ) } \end{gathered}$ | Quantity <br> 1×96 well kit | Reosaturtion |
| :---: | :---: | :---: |
| Nuclease ELISA plate <br> Stripwell microplate with 96 anti-nuclease <br> monocional antibodies coated wells | $\begin{gathered} 96 \text { wells } \\ 12 \times 8 \text { wel stips }) \end{gathered}$ | Ready for use |
| Standard <br> Nuclease lyophilized from buffered protein solution with preservatives | $\stackrel{1 \text { vial }}{\text { ayophilize torm) }}$ | Refer to the vial label for reconstitution volume. Reconstitute by adding Standard reconstitution buffer to be a stock solution of $50 \mathrm{ng} / \mathrm{mL}$. (see procedure, section 8.(2)) |
| Standard reconstitution buffer Buffered protein solution with preservatives | $\begin{gathered} 1 \text { Val } \\ \text { (1.5 mu) } \end{gathered}$ | Ready for use |
| HRP-antibody conjugate <br> HRP conjugated ant-nuclease moniocional antibody in buffered protein solution with preservatives | $\begin{aligned} & \begin{array}{l} \text { 1 vial } \\ (0, \mu \mathrm{~L}) \end{array} \end{aligned}$ | Dilute $200 \times$ with HRP-antibody conjugated dilluent buffer (see reagent preparation, section 5A. |
| HRP-antibody conjugated diluent buffer Buffered solution with preservatives | $\begin{gathered} \text { 1 Val } \\ (12 \mathrm{~mL}) \end{gathered}$ | Ready for use |
| 20 X wash buffer <br> 20 -fold concentrated solution of buffered surfactant with preservatives | $\begin{gathered} \text { 1 val } \\ 15 \mathrm{~mm} \end{gathered}$ | Dilute $20 \times$ with distilled water (see reagent preparation section 5.B) |
| TMB <br> Chromogenic substrate (tetramethylbenzidine) for HRP | $\begin{gathered} 1 \text { val } \\ \text { (12 mu) } \end{gathered}$ | Ready for use |
| Stop solution $\mathrm{H}_{2} \mathrm{SO}_{4}$ solution | $\begin{gathered} 1 \text { vial } \\ (6 \mathrm{~mL}) \end{gathered}$ | Reedy toru |
| Microplate sealing film | 1 sheet | N/A |

Reagents provided and reconstitution

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## Taipei Office

