

| Product Information |                |  |
|---------------------|----------------|--|
| PerfeСта DNAse I    |                |  |
| Part Number         | 95150-01K      |  |
| Number of Reactions | 1000 Reactions |  |
| Concentration       | 2 units/μL     |  |
| Storage Temperature | -25°C to -15°C |  |
| Lot Number          | 022995         |  |
|                     | 014988         |  |
| Reference Number    | 061716         |  |
|                     | 011117         |  |
| Expiration Date     | 06/30/19       |  |

## **Product Description:**

PerfeCTa DNase I is an ultra-pure, recombinant bovine DNase I preparation that is permanently inactivated with a proprietary stop buffer and simple heat kill incubation. This reagent provides a simple and rapid solution to eliminate residual genomic DNA from total RNA preparations for expression profiling by reverse transcription quantitative PCR amplification (RT-qPCR) as well as other molecular biology applications. Complete and permanent DNase I inactivation is critical for linear first-strand cDNA synthesis. Residual or renatured DNase will degrade cDNA product and lead to inaccurate target quantification. One unit completely degrades 1µg of dsDNA in 10 minutes at 37°C.

## Component Part Numbers:# 84271 PerfeCTa DNase I 0.500 mL

| Product Specifications |                                      |                                 |       |
|------------------------|--------------------------------------|---------------------------------|-------|
| 95150                  |                                      |                                 |       |
| Assay                  | qRT-PCR Functional Assay for DNAse I | DNasel Mu-gDNA Functional Assay | RNase |
| Result                 | Pass                                 | Pass                            | Pass  |

## **Quality Control Analysis and Specifications:**

**RNase Contamination:** RNase activity must be below the detectable limits of 1 pg RNase A equivalent as assayed using a fluorogenic substrate following a 1 hour incubation at 37°C with each kit component at 1X concentration.

**qRT-PCR Functional Assay for DNAse I:** The Ct difference between positive (+) DNase and Negative (-) DNase I samples must be  $\leq$  1.5 Ct

**DNasel Mu-gDNA Functional Assay:** The Ct difference between positive (+) DNase and Negative (-) DNase I samples must be  $\geq 15$  Ct

## **Limitations of Use**

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