

Instructions for Use

Life Science Kits & Assays



innuDETECT Clostridium perfringens Assay

1 Product specifications

| Starting material | Isolated total DNA |
|-------------------------|--|
| Time of detection | ~ 90 minutes |
| qPCR detection channels | FAM (<i>Clostridium perfringens</i>) HEX (Internal Control) |

The Assay detects DNA derived from sample material using an extraction kit suitable to isolate bacterial DNA. Please make sure that the common quality requirements for DNA samples are achieved.

2 Intended use

The innuDETECT *Clostridium perfringens* Assay has been designed to detect the DNA derived from *Clostridium perfringens* using TaqMan principle.

Clostridium perfringens is a rod-shaped, gram-positive, endospore-forming, flagellate bacterium.

Clostridium perfringens triggers gastroenteritis after absorption into the body, the symptoms begin 6-24 h after ingestion of contaminated food or water. In order to be able to detect parasitic contamination within the framework of drinking water quality monitoring, waters originating from or influenced by surface water must be routinely tested for *Clostridium perfringens*.

The assay detects the *Clostridium perfringens* specific *cpa*-gene (detected in FAM-channel).

The assay includes an Internal Control (IC) that can be used as amplification control if added to the PCR reaction. If added to

the Lysis Buffer the IC can also be utilized to check the used DNA extraction method.

The assay is intended for research use only.

3 Product and order number

| Name | Amount | Order-no. |
|--|--------|-----------------|
| innuDETECT Clostridium perfringens Assay | 24 rxn | 845-IDF-0034024 |
| innuDETECT Clostridium perfringens Assay | 96 rxn | 845-IDF-0034096 |






4 Storage conditions

The Assay is delivered at ambient temperature.

The kit is shipped at ambient temperature. Upon arrival store the innuDETECT Clostridium perfringens Assay at -22 °C to 18 °C, except the innuDRY qPCR MasterMix Probe that before dissolving should be stored at 4 °C – 8 °C.

When stored as recommended, the kit is stable until the expiration date printed on the label on the kit box.

5 Delivered components

| Components | | ∑ 24 | ∑ 96 |
|--|---|------------|-------------|
| Primer/Probe Mix Clostridium perfringens/IC |  | 75 µl | 300 µl |
| innuDRY qPCR MasterMix Probe |  | 1 (25 rxn) | 1 (100 rxn) |
| Resuspension Buffer Probe |  | 300 µl | 1.1 ml |
| Internal Control |  | 1 | 1 |
| PCR-grade H ₂ O |  | 2 ml | 2 x 2 ml |

6 Safety precautions

The assay shall only be handled by educated personal in a laboratory environment. The compliance with the specified procedure is absolute mandatory when performing this assay.

Reagents should be stored in their original containers at the indicated temperatures. Do not replace individual components with those from different batches or test assays. Note the indicated expiration dates.

Do not eat, drink or smoke while performing the assay.

Wear protective clothing and safety gloves.

All samples and test materials should be handled and disposed of as infectious material, in accordance with regulatory requirements.

Reagent containers that have not come in contact with potentially infectious material may be disposed of along with ordinary laboratory waste.

Store the reagents used for performing PCR separately from DNA templates and amplification products.

7 Reagent preparation

7.1 Internal Control

Dissolve the lyophilized Internal Control (IC) by adding 1.25 ml of PCR-grade H₂O and mixing thoroughly.

To use the IC as an amplification control, add the IC to the qPCR Reaction Mix in an amount of 0,1 µl per reaction. NOTE: the No Template Control (NTC) must also be positive for IC

To use the IC as an extraction control, add to the Lysis Buffer/Sample Mix an amount of IC which is 1/10 of the final elution volume (see corresponding DNA isolation manual). To observe the relative loss of DNA during the extraction procedure, perform the amplification of 1µl of 1:10 diluted IC added to the NTC. This corresponds to 100% DNA extraction efficiency.

7.2 2x qPCR MasterMix Probe

The 2x innuDRY qPCR MasterMix Probe must be prepared before starting the PCR setup and can be stored at -22 °C to -18 °C.

Add 250 µl (for 24 rxn assay) or 1 ml (for 96 rxn assay) of Re-suspension Buffer Probe to the innuDRY qPCR MasterMix Probe tube. Incubate the tube in a thermal shaker for 20 minutes at 25°C and 550 rpm. Spin down the tube to collect the liquid on the bottom.

8 Real-Time PCR (qPCR)

8.1 Preparation of Reaction Mix

| Reagent | Volume (1 rxn) |
|--|-------------------------------------|
| 2x qPCR MasterMix Probe | 10 μ l |
| Primer/Probe Mix Clostridium perfringens/IC | 3 μ l |
| IC, diluted 1:10 (if used as amplification IC) | 1 μ l |
| Sample (PCR-grade H ₂ O for NTC) | \leq 2 μ g DNA, max 5 μ l |
| PCR-grade H ₂ O | Fill up to 20 μ l |

Determine the total number of required qPCR reactions considering your Samples and at least one NTC.

The composition of the qPCR Reaction Mix per single reaction (1 rxn) is shown in the table below. Prepare the qPCR Reaction Mix for the number of reactions needed with a 10 % surplus volume to allow for pipetting error.

To avoid the cross contamination first aliquot the qPCR Reaction Mix to PCR stripe or plate and then add samples to the qPCR reaction mix.

8.2 Real-Time PCR Conditions

For basic information regarding the setup and programming of the different Real-Time PCR device, please refer to the manual of the respective instrument.

Program the Real-Time PCR device as indicated in the table below and start the program.

Interpretation of results

| Step | Cycles | Profile | Temperature | Retention time |
|------|--------|-------------------------|-------------|----------------|
| 1 | 1 | Initial Denaturation | 95 °C | 120 s |
| 2 | 45 | Denaturation | 95 °C | 15 s |
| | | Annealing / Elongation* | 55 °C | 60 s |

* Data acquisition: Fluorescence Detection (FAM; HEX)

9 Interpretation of results

Please refer to the following table to identify the signal pattern that matches to the obtained signals. It is strongly recommended to run at least one No Template Control (NTC) for each experiment.

The Ct value of IC can vary (or even disappear) in dependence of DNA quality and intensity of FAM signal.

If the sample is strongly positive for *Clostridium perfringens*, in rare cases it is possible that the IC is negative.

| FAM | HEX | Sample | Valid | Recommended interpretation |
|-----|-----|----------------|-------|--|
| + | + | NTC | no | Contamination of PCR chemicals with target or (and) IC DNA |
| - | + | NTC | yes | No contamination |
| + | + | Unknown Sample | yes | Positive for <i>Clostridium perfringens</i> |
| - | + | Unknown Sample | yes | Negative for <i>Clostridium perfringens</i> |
| - | - | Unknown Sample | no | PCR reaction and/or DNA isolation failed |
| + | - | Unknown Sample | yes | The sample is strong positive for the target |

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