Instructions for Use Life Science Kits & Assays



innuDETECT Pork Assay



1 Product specifications

Starting material	Isolated total DNA
Time of detection	~ 60 minutes
qPCR detection channels	FAM (Target) and HEX (IC)
Sensitivity	Up to 5 DNA copies/PCR

Detection of DNA isolated from sample material using an extraction kit suitable to isolate total DNA. A DNA isolation kit from IST Innuscreen GmbH is highly recommended (see Related Products). Please make sure that the common quality requirements for DNA samples are achieved.

2 Intended use

The innuDETECT Pork Assay is a molecular diagnostic test system for detection of pork DNA in different samples based on TaqMan[®] principle.

The target sequence is a mitochondrial gene (Cytochrome b). Therefore, even very small amounts of target DNA (up to 1 pg per PCR-reaction) can lead to positive results.

The assay includes a Positive Control (PC) with 10.000 target copies/µl.

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 DNAextraction method used.

The assay is intended for research use only.

3 Product and order number

Name	Amount	Order-no.
innuDETECT Pork Assay	24 rxn	845-IDF-0010024
innuDETECT Pork Assay	96 rxn	845-IDF-0010096

4 Storage conditions

The Assay is delivered at ambient temperature.

Store the innuDETECT Pork Assay at -22 °C to -18 °C, except the innuDRY qPCR MasterMix Probe that should be stored before dissolving at 4 °C to 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 Pork IC	75 µl	300 µl
innuDRY qPCR MasterMix Probe	1	1
Resuspension Buffer Probe	300 µl	1.1 ml
Positive Control	30 µl	30 µl
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 absolutely 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 H_2O and mix thoroughly.

To use the IC as an <u>amplification control</u>, add 1μ I of IC to each PCR reaction.

Alternatively, the IC can be added to the qPCR reaction mix in an amount of 1 μ l/reaction. <u>NOTE</u>: in this case the No Template Control (NTC) must also be positive for IC.

To improve the sensitivity of the test for samples with a very low target amount, reduce the IC content up to 1:10.

To use the IC as an <u>extraction control</u>, add to the Lysis Buffer/Sample Mix amount of IC which is 1/10 from final elution volume (see according DNA isolation instruction manual). Use the co-amplification of spiked IC to observe the relative loss of DNA during the extraction procedure.

7.2 2x MasterMix

The 2x MasterMix 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 Resuspension Buffer Probe to the innuDRY qPCR MasterMix Probe tube. Vortex carefully and centrifuge the tube to collect the liquid on the bottom.

8 Real-Time PCR (qPCR)

8.1 Preparation of reaction batches

Determine the total number of required qPCR reactions considering also at least one NTC.

The composition of the qPCR reaction mix for one sample is shown in the table below. Prepare the qPCR reaction mix for the number of samples needed (including NTC).

Add qPCR reaction mix to PCR stripe or plate. In a second step add samples to the qPCR reaction mix in order to avoid the cross contamination.

Reagent	Volume (1 rxn)		
2x MasterMix	10 µl		
Primer/Probe Mix Pork IC	3 µl		
IC (if used as amplification IC)	1 µl		
Sample (PCR-grade H ₂ O for NTC)	≤ 2 µg DNA, max 5 µl		
or PC (if needed)	1 µl		
PCR-grade H ₂ O	Fill up to 20 µl		
Seal the PCR stripe or plate with an appropriate sealing film			
(PP) and/or cap; place tubes in the Real-Time PCR Cycler and			
close the lid.			

8.2 Real-Time PCR conditions

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

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

Step	Cycles	Profile	Temperature	Retention time
1	1	Initial denaturation	95 °C	120 s
2		Denaturation	95 ℃	10 s
	35	Annealing / Elongation*	62 °C	45 s
4 D .				

* 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.

FAM	HEX	Sample	Valid	Recommended interpretation
+	(+)	NTC	no	Contamination of PCR chemicals with target or (and) IC DNA
-	+	NTC	yes	No contamination
+	(+)	PC	yes	PCR run successful
+/-	+	Unknown Sample	yes	Positive or negative for the target
-	-	Unknown Sample	no	PCR reaction and/or DNA isolation failed
++	-	Unknown Sample	yes	The sample is strong positive for the target

10 Related products

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