Instructions for Use Life Science Kits & Assays





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 Analytik Jena is highly recommended (see Related Products). Please make sure that the common quality requirements for DNA samples are achieved.

2 Intended use

The innuDETECT Sheep Assay is a molecular diagnostic test system for detection of sheep 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 DNA-extraction method used.

The assay is intended for research use only.

3 Product and order number

Name	Amount	Order-no.
innuDETECT Sheep Assay	24 rxn	845-IDF-0050024
innuDETECT Sheep Assay	96 rxn	845-IDF-0050096

4 Storage conditions

Store the innuDETECT Sheep 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 Sheep 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 H₂O	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 1 µl 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. NOTE: 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 μΙ
Primer/Probe Mix Sheep IC	3 µl
IC (if used as amplification IC)	1 μΙ
Sample (PCR-grade H₂O for NTC)	≤ 2 µg DNA, max 5 µl
or PC (if needed)	1 μΙ
PCR-grade H₂O	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 ℃	120 s
2		Denaturation	95 ℃	10 s
	40	Annealing / Elongation*	62 °C	45 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.

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

Order Number
845-KS-1040
845-KS-1540
845-IPP-5716
845-BP-0010
845-IDF-0010
845-IDF-0020
845-IDF-0030
845-IDF-0040
845-IDF-0060
845-IDF-0070
845-IDF-0080
845-IDF-0090
845-IDF-0100
845-IDF-0110
845-IDF-0120
845-IDF-0130

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