



Instructions for life science research use only. Not tested for use in diagnostic procedures. For *in vitro* use only.



## Instructions For Use

# LightMix<sup>®</sup> Modular MSTN Extraction Control

580

Cat.-No. 58-0905-96

Roche SAP n° 07 559 968 001

Kit with reagents for 96 PCR reactions 20 µl for detection of MSTN [lyophilized]

## 1. Content, Storage and Expiry

- 1 Vial red cap 96 reactions MSTN (lyophilized)
- 1 Vial seek dye ROX (330 µl)

## Storage at Arrival:

Store cooled or at ambient temperature  
Do **not** freeze the lyophilized reagents.

- Lyophilized kits are stable for at least 6 months (4°C to 25°C in the dark). See lot-specific expiry date.
- Dissolved reagents are stable for at least 2 weeks if stored protected from light and cooled (4°C).
- Dissolved reagents can be stored long-term at -20°C (within expiry). Avoid multiple freeze-thaw cycles.

## 2. Additional Reagents required

LightCycler<sup>®</sup> Multiplex RNA Virus Master  
LightCycler<sup>®</sup> Multiplex DNA Master  
or Roche LightCycler<sup>®</sup> 480 Probes Master (no instructions included)

Cat.-No. 06 754 155 001

Cat.-No. 07 339 585 001

Cat.-No. 04 707 494 001

## 3. Introduction

PCR analysis of biological samples may occasionally result difficult due to problems related to the extraction process. In particular, false negative results in pathogen testing can be due to the presence of inhibitors, degraded target material or an unsuccessful extraction of nucleic acids. The presence of amplifiable nucleic acids can be verified by running an extraction control PCR.

This product is intended to be used as extraction control and as reference gene for DNA or RNA-based assays. The targeted region of the Myostatin (MSTN) gene is highly conserved in chordata.

For use with capillary LightCycler instruments include a seek dye to ensure to detect the capillaries.

## 4. Description

A 89 bp long fragment from the myostatin gene or transcript is amplified with specific primers and detected with a hydrolysis probe with a R6G labeled hydrolysis probe. Starting with lot 3407 this kit has been improved by adding one primer and one probe to detect a wider variety of species.

## 5. Specification

These reagents detect 0.01 ng or less of *vertebrate* DNA. The linear measuring range is 0.01 to 500 ng DNA.

The product has been confirmed to work with samples from human (*Homo sapiens*), porc (*Sus suis*), beef (*bos bovis*), dromedary (*Camelus dromedarius*), chicken (*Gallus gallus*), turkey (*Meleagris gallopavo*) ostrich (*Struthio camelus*), mallard (*Anas platyrhynchos*), pigeon (*Columba livia*), quail (*Coturnix coturnix*) guinea fowl (*Numida meleagris*), duck muscovy (*Cairina moschata*), and goose (*Anser ansermallard*).

## 6. Sample Material and Extraction

Depends on the analytical PCR (DNA or RNA). See ModularDx Document **Extraction Protocols**.

## 7. Material Safety Data (MSDS)

According to OSHA 29CFR1910.1200, Australia [NOHSC:1005, 1008 (1999)] and the EU Directives 67/548/EC and 1999/45/EC any products which do not contain more than 1% of a component classified as hazardous or classified as carcinogenic do not require a Material Safety Data Sheet (MSDS).

Product is not hazardous, not toxic, not IATA-restricted. Product is not from human, animal or plant origin. Product contains synthetic oligonucleotide primers and probes.



## 8. Instructions for Use

- Instrument programming see document **ModularDx Programming**
- Color Compensation see instructions in **40-0320 Universal Color Compensation Hexaplex**
- Pipetting instructions multiplex PCR see **ModularDx Multiplex**

### 8.1. Programming Roche 480 Instruments

See the Instrument operator's manual for details. Start programming before preparing the solutions. The protocol consists of four program steps:

- 1: Reverse Transcription of the viral RNA
- 2: Denaturation: sample denaturation and enzyme activation
- 3: Cycling: PCR-amplification
- 4: Cooling: cooling the instrument

Detection Format 580 Channel	Set Quant Factor 10, Max Integration time 1 sec
LightCycler® 480 Instrument:	523-568
LightCycler® 480 II Instrument:	533-580
cobas z 480 Analyzer (open channel):	540-580

Program Step:	RT Step	Denaturation	Cycling			Cooling
<b>Parameter</b>						
Analysis Mode	None	None	Quantification mode			None
Cycles	1	1	45			1
Target [°C]	55	95	95	60	72	40
Hold [hh:mm:ss]	00:05:00	00:05:00	00:00:05	00:00:15	00:00:15	00:00:30
Ramp Rate [°C/s] 96	4.4	4.4	4.4	2.2	4.4	1.5
Ramp Rate [°C/s] 384	4.6	4.6	4.6	2.4	4.6	2.0
Acquisition Mode	None	None	None	Single	None	None

\* 1-Step RT-PCR optional

Table 1

### 8.2. Experimental Protocol

- **Sample material:** Use aqueous nucleic acid preparations (e.g. 'High Pure Viral Nucleic Acid Kit').
- **Negative control:** Always run at least one no-template control (NTC) - replace the template NA with water.

For an increased sensitivity use 10 µl sample per 20 µl reaction, in case that inhibition is likely to occur, e.g. extracts obtained from fecal samples, use 5 µl. For 10 µl reactions in 384 well plates use 5 µl /2.5 µl.

#### 8.2.1. Preparation of Parameter-Specific Reagents (PSR, 96 reactions):

One reagent vial with a **red** cap contains all primers and probe to run 96+ LightCycler® reactions.

**Add 50 µl** PCR-grade water to each reagent vial, mix the solution (vortex) and spin down. For robotic pipetting the volume can be extended to 55 µl (signals will decrease by 10-20%).

► Use 0.5 µl reagent for a 20 µl PCR reaction.

#### 8.2.2. Preparation of the Target Nucleic Acids

MSTN is a control reaction working with DNA and RNA. Follow the procedure for the analytical PCR.

### 8.2.3. Preparation of the Reaction Mix RNA/DNA Multiplex Master

Duplex PCR - Instruction for use with the Roche LightCycler® Multiplex RNA Virus Master		
for 5 µl extract	Component	10 µl extract
9.9 µl	Water, PCR-grade (colorless cap, provided with the Roche Master kit)	4.9 µl
0.5 µl	<b>MSTN Control Reaction and</b>	0.5 µl
<b>0.5 µl</b>	<b>Reagent mix additional assay(s) (Multiplex PCR))</b>	<b>0.5 µl</b>
--	LightCycler® 1.x / 2.0 optional : 3 µl Seek dye (substitute water)	--
4.0 µl	<b>Roche Master</b> (see Roche manual)	4.0 µl
0.1 µl	<b>RT Enzyme</b> (RNA Master only)	0.1 µl
<b>15.0 µl</b>	<b>Volume of Reaction Mix (DNA Master 14.9 µl)</b>	<b>10.0 µl</b>

Table 2

Mix gently, spin down and **transfer 15 µl (10 µl)** per well.

**Add 5 µl (10 µl)** of sample or control to each well for a final reaction volume of 20 µl. Seal plate and centrifuge.

**Start run**

### 9. Typical Results (Data from LightCycler® 480 II system)

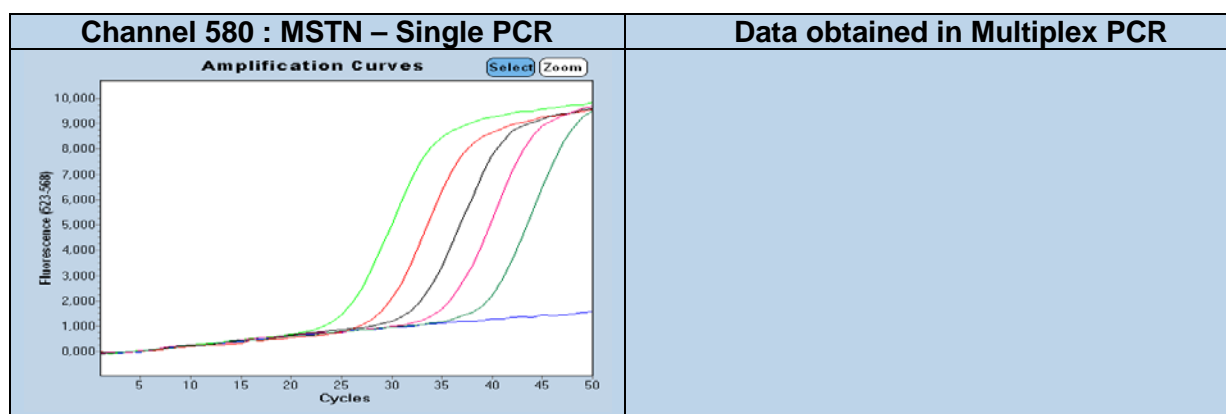


Figure 1

### 10. Reading the Results

Perform data analysis as described in the operator's manual. For multiplex assays select the color compensation. We recommend using the Second Derivative Maximum method (Automated (F" max). View results in the 580 channel. The negative control (NTC) and negative samples must show a signal.

Channel 580 Control Reaction	Other channel Analytical PCR	Analytical PCR NTC Control	Result
Amplification +	Not detectable	Negative	Parameter - Negative
Not relevant	Amplification signal	Negative	Parameter- Positive
No amplification	Not detectable	Negative	PCR failure Repeat
Not relevant	Not relevant	Positive	Contamination Repeat

**Notes:** cobas z 480 Analyzer signal levels are about 50% compared to LightCycler® 480 II results.  
+ Cp depends on the respective dilution during extraction.

### 11. References

Analytical Performance Determination and Clinical Validation of the Novel Roche RealTime Ready Influenza A/H1N1 Detection Set. Wenzel et al. (2010)

## 12. Multiplex PCR Compatibility

The MSTN extraction control assay can be combined with up to three (for LightCycler® 96 instruments), four (cobas z 480 analyzer) or five (LightCycler® 480 systems) analytical assays :

**Multiplex PCR and Instrument Compatibility**  
Color Compensation 40-0320 is mandatory for Multiplex PCR


500	530	580	610	640	660
	Assay 1	MSTN			
	Assay 1	MSTN	Assay 2		Assay 3
	Assay 1	MSTN	Assay 2	Assay 4	Assay 3
Assay 5	Assay 1	MSTN	Assay 2	Assay 4	Assay 3

480 II	z 480	LC96	LC2.0	Nano
X	X	X	X	X
X	X	X		
X	X			
X				

Table 3

## 13. Version History

V140404	Release version	2014-04-14
V140909	Editorial changes	2014-09-09
V150101	Editorial changes	2015-02-27
V150525	Roche SAP number changed Sample volume, instrument settings and PCR cycles (Multiplex Master) <b>One probe added</b> - spectrum of detected (bird) species enlarged	2015-05-25
V151001	2015 protocol Multiplex Master, 5/10 µl extract and 60°C acquisition	2015-10-01
V160313	8.2.3 wording	2016-07-07

<b>Certificate of Analysis (CoA)</b>						
Lot n°						
Expiry :						
<b>Dilution</b>	<b>Placenta</b>	<b>DNA</b>	<b>100 ng</b>	<b>10 ng</b>	<b>1 ng</b>	<b>passed</b>
<b>Cp range</b>			23-26	26-29	30-33	
<b>Measured</b>						
<b>Signal level</b>			20-40			
<b>Measured</b>						
<b>Negatives</b>	<b>10/10</b>					
<p><b>Note:</b> Cp (crossing point) values collected with pDNA (single target PCR). Fluorescence (FL) levels depend on instrument settings and may vary. The Cp values will vary from instrument to instrument by up to 2 cycles, while the distance between two dilution steps should be relative constant (<math>\Delta</math>Cp).</p>						
QC Acceptance Date:			YYYYMMDD			
We, the undersigned, certify that the product designated above has been obtained in accordance with the rules of production and quality control.						
<b>Name(s) :</b>						

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