

## Instructions for use

# FlexStar<sup>®</sup> alpha-CoV & beta-CoV & RV RT-PCR Detection Mix 1.5

01/2024 EN

Respiratory

**FlexStar**<sup>®</sup>

# **FlexStar**<sup>®</sup>

# alpha-CoV & beta-CoV & RV RT-PCR Detection Mix 1.5

For research use only!

(RUO)



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## 1. Application

The FlexStar<sup>®</sup> alpha-CoV & beta-CoV & RV RT-PCR Detection Mix 1.5 is a reagent system, based on real-time PCR technology, for the qualitative detection and differentiation of RNA from the alpha coronaviruses (CoV) human CoV-229E and human CoV-NL63, RNA from the beta coronaviruses human CoV-OC43 and human CoV-HKU1 and rhinovirus (RV) specific RNA.

For research use only (RUO)! Not for use in diagnostic procedures.

## 2. Product content

The FlexStar<sup>®</sup> alpha-CoV & beta-CoV & RV RT-PCR Detection Mix 1.5 contains the following components:

Lid color	Component	Number of tubee	Nominal volume [µl/tube]
Blue	Detection Mix <sup>1)</sup>	8	60
Red	PC <sup>2)</sup>	2	250
White	NTC <sup>3)</sup>	2	250

#### Table 1: Kit components

<sup>1)</sup> Contains biological material of animal origin

<sup>2)</sup> Positive Control (alpha-CoV, beta-CoV and RV specific RNA)

<sup>3)</sup> No Template Control (negative control)

## 3. Storage

- The FlexStar<sup>®</sup> alpha-CoV & beta-CoV & RV RT-PCR Detection Mix 1.5 is shipped on dry ice. The product components should arrive frozen. If one or more components are not frozen upon receipt, or if tubes have been compromised during shipment, contact altona Diagnostics technical support for assistance (see chapter 9. Technical assistance).
- All components should be stored at -25 °C to -15 °C upon arrival.
- Repeated thawing and freezing of the Detection Mix component should be avoided, as this might affect the performance of the product.
- Repeated thawing and freezing of the Positive Control (PC) and No Template Control (NTC) (more than 4 times) should be avoided, as this might affect the performance of the product.
- Storage at room temperature (max. +30 °C) should not exceed a period of 2 hours.
- Protect the Detection Mix component from light.

## 4. Product description

The FlexStar<sup>®</sup> alpha-CoV & beta-CoV & RV RT-PCR Detection Mix 1.5 is a reagent system. Used in combination with the FlexStar<sup>®</sup> (RT-)PCR Amplification Mix 1.5 it allows the qualitative detection and differentiation of NL63/229E, OC43/HKU1 and RV specific RNA.

The FlexStar<sup>®</sup> alpha-CoV & beta-CoV & RV RT-PCR Detection Mix 1.5 is based on real-time RT-PCR technology, utilizing reverse-transcriptase (RT) reaction to convert RNA into complementary DNA (cDNA), polymerase chain reaction (PCR) for the amplification of NL63/229E, OC43/HKU1 and RV specific target sequences and fluorescently labeled target specific probes for the detection of the amplified cDNA. In addition to the NL63/229E, OC43/HKU1 and RV RNA specific amplification and detection systems, the FlexStar<sup>®</sup> alpha-CoV & beta-CoV & RV RT-PCR Detection Mix 1.5 includes oligonucleotides for the amplification and detection of the internal control (IC, AltoStar<sup>®</sup> Internal Control 1.5). The IC is automatically added at the beginning of the nucleic acid purification procedure on the AltoStar<sup>®</sup> Automation System AM16 (in the following summarized as AltoStar<sup>®</sup> AM16). For details refer to the instructions for use of the AltoStar<sup>®</sup> Internal Control 1.5.

Probes specific for NL63/229E RNA are labeled with the fluorophore FAM<sup>™</sup>, probes specific for OC43/HKU1 RNA are labeled with the fluorophore ROX<sup>™</sup> and probes specific for RV RNA are labeled with the fluorophore Cy5, respectively. The probe specific for the IC is labeled with the fluorophore JOE<sup>™</sup>.

Using probes linked to distinguishable dyes enables the parallel detection of NL63/229E, OC43/HKU1, RV and the IC in the corresponding detection channels of the real-time PCR instrument.

## 4.1 Components

The FlexStar<sup>®</sup> alpha-CoV & beta-CoV & RV RT-PCR Detection Mix 1.5 contains enough reagents for 96 reactions. The product consists of the following components:

- Detection Mix<sup>1)</sup>
- PC<sup>2)</sup>
- NTC<sup>3)</sup>
- <sup>1)</sup> Contains biological material of animal origin
- <sup>2)</sup> Positive Control (alpha-CoV, beta-CoV and RV specific RNA)
- <sup>3)</sup> No Template Control (negative control)

Except for the DNA polymerase and the reverse transcriptase, which are included in the FlexStar<sup>®</sup> (RT-)PCR Amplification Mix 1.5, the Detection Mix component contains all reagents (PCR buffer, magnesium salt, primers and probes) to allow detection and differentiation of NL63/229E and OC43/HKU1 and detection of RV specific RNA, as well as of IC specific RNA.

The PC contains alpha-CoV and beta-CoV as well as RV specific RNA. It is used to verify the functionality of the NL63/229E, OC43/HKU1 and RV RNA specific amplification and detection systems.

The NTC contains neither NL63/229E, OC43/HKU1, nor RV specific RNA but does contain the IC template. The NTC is used as negative control for the NL63/229E, the OC43/HKU1 and the RV RNA specific real-time PCR and indicates possible contamination of the Detection Mix component.

## 4.2 Real-time PCR instruments

The FlexStar<sup>®</sup> alpha-CoV & beta-CoV & RV RT-PCR Detection Mix 1.5 can be used with the following real-time PCR instruments:

- CFX96™ Deep Well Dx System (Bio-Rad)
- CFX96<sup>™</sup> Dx System (Bio-Rad)
- QuantStudio<sup>™</sup> 5 Real-Time PCR System (Applied Biosystems)
- Rotor-Gene® Q5/6 plex Platform (QIAGEN)

#### NOTE

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Ensure that all instruments used have been installed, calibrated, checked and maintained according to the manufacturer's instructions and recommendations.

## 5. Material required but not provided

The following additional instruments and consumables are required for use of the FlexStar<sup>®</sup> alpha-CoV & beta-CoV & RV RT-PCR Detection Mix 1.5 but not provided with this product:

- Appropriate real-time PCR instrument (see chapter 4.2 Real-time PCR instruments)
- Appropriate nucleic acid extraction system or kit (see chapter 6.1 Sample preparation)

- Vortex mixer
- · Centrifuge (e.g., desktop centrifuge) for centrifugation of kit reagents
- Centrifuge for centrifugation of PCR plates
- Appropriate 96 well reaction plates or reaction tubes with corresponding (optical) closing material
- Pipettes (adjustable)
- Pipette tips with filters (disposable)
- Powder-free gloves (disposable)

Reagents required but not included in the FlexStar<sup>®</sup> alpha-CoV & beta-CoV & RV RT-PCR Detection Mix 1.5:

- FlexStar® (RT-)PCR Amplification Mix 1.5 (Order No. FS0011503/FS0011505)
- AltoStar<sup>®</sup> Internal Control 1.5 (Order No. IC15-06)

## 6. Procedure

### 6.1 Sample preparation

Extracted RNA is the starting material for the FlexStar<sup>®</sup> alpha-CoV & beta-CoV & RV RT-PCR Detection Mix 1.5. The quality of the extracted RNA has a profound impact on the performance of the product.

For additional information and technical support regarding pre-treatment and sample preparation, contact altona Diagnostics technical support (see chapter 9. Technical assistance).

#### 6.2 Master mix setup

All reagents and samples should be thawed completely, mixed (by pipetting or gentle vortexing) and centrifuged briefly before use.

The FlexStar<sup>®</sup> alpha-CoV & beta-CoV & RV RT-PCR Detection Mix 1.5 is configured for use with the FlexStar<sup>®</sup> (RT-)PCR Amplification Mix 1.5 and the AltoStar<sup>®</sup> Internal Control 1.5, which allows to control the sample preparation procedure (nucleic acid extraction) and the subsequent RT-PCR.

- ► The IC is automatically added at the beginning of the nucleic acid purification procedure on the AltoStar<sup>®</sup> AM16.
- When using other nucleic acid extraction methods, the IC has to be added during the lysis step either manually or automatically by the respective instrument.
- No matter which method/system is used for nucleic acid extraction, never add the IC directly to the sample. The IC should always be added to the sample/ lysis buffer mixture. The volume of the IC which has to be added, always and only depends on the elution volume. It represents 50 % of the elution volume. For instance, if the nucleic acid is going to be eluted in 60 µl of elution buffer or water, 30 µl of IC per sample must be added into the sample/lysis buffer mixture.
- Set up the master mix according to the following pipetting scheme:

Number of reactions (rxns)	1	12
Detection Mix	5 µl	60 µl
Amplification Mix	15 µl	180 µl
Volume master mix	20 µl	240 µl

 Table
 2: Pipetting scheme (master mix setup)

#### NOTE

No matter which method/system is used for nucleic acid extraction, never add the IC directly to the specimen.

#### 6.3 Reaction setup

- Pipette 20 µl of the master mix into each required well of an appropriate optical 96-well reaction plate or an appropriate optical reaction tube.
- Add 10 µl of the sample (eluate from the nucleic acid extraction) or 10 µl of the controls (PC or NTC).

Table 3: Pipetting scheme (reaction setup)

Reactio	n setup
Master mix	20 µl
Sample or control	10 µl
Total volume	30 µl

- ▶ Make sure that at least 1 PC and 1 NTC is used per run.
- Thoroughly mix the samples and controls with the master mix by pipetting up and down.
- Close the 96-well reaction plate with appropriate lids or optical adhesive film and the reaction tubes with appropriate lids.
- Centrifuge the 96-well reaction plate in a centrifuge with a microtiter plate rotor and the reaction tubes in an appropriate centrifuge for 30 seconds at approximately 1,000 x g (~ 3,000 rpm).
- ► The NTC does already contain the IC template in the correct concentration.

## 7. Programming the real-time PCR instrument

For basic information regarding the setup and programming of the different realtime PCR instruments, refer to the user manual of the respective instrument.

For detailed programming instructions regarding the use of the FlexStar<sup>®</sup> alpha-CoV & beta-CoV & RV RT-PCR Detection Mix 1.5 on specific real-time PCR instruments, contact altona Diagnostics technical support (see chapter 9. Technical assistance).

## 7.1 Settings

► Define the following settings:

#### Table 4: Run settings

Settings		
Reaction volume	30 µl	
Ramp rate	Default	
Passive reference*	None	

\* If applicable

## 7.2 Fluorescence detectors (dyes)

Define the fluorescence detectors (dyes):

#### Table 5: Fluorescence detectors

Target	Detector name	Reporter	Quencher
HCoV-229E and HCoV-NL63 specific RNA	NL63/229E	FAM™	(None)
HCoV-OC43 and HCoV-HKU1 specific RNA	OC43/HKU1	ROX™	(None)
Rhinovirus specific RNA	Rhinovirus	Cy5	(None)
IC	Internal Control	JOE™	(None)

## 7.3 Temperature profile and dye acquisition

▶ Define the temperature profile and dye acquisition:

Table 6: Temperature profile and dye acquisition

	Stage	Cycle repeats	Acquisition	Temperature [°C]	Time [min:s]
Reverse transcription	Hold	1	-	52	05:00
Denaturation	Hold	1	-	95	00:05
Amplification	plification Cycling	45	-	95	00:05
Amplification			Yes	58	00:25

## 8. Data analysis

For basic information regarding data analysis on specific real-time PCR instruments, refer to the user manual of the respective instrument.

For detailed instructions regarding the analysis of the data generated with the FlexStar<sup>®</sup> alpha-CoV & beta-CoV & RV RT-PCR Detection Mix 1.5 on different realtime PCR instruments, contact altona Diagnostics technical support (see chapter 9. Technical assistance).

## 8.1 Interpretation of results

#### 8.1.1 Qualitative analysis

#### Table 7: Result interpretation

Detection channel				
FAM™ (NL63/ 229E)	ROX™ (OC43/ HKU1)	Cy5 (Rhino- virus)	JOE™ (IC)	Result interpretation
+	+	+**	+/-*	NL63/229E, OC43/HKU1 and RV specific RNA detected.
+	+	-	+/-*	NL63/229E and OC43/HKU1 specific RNA detected.
+	-	+**	+/-*	NL63/229E and RV specific RNA detected.
-	+	+**	+/-*	OC43/HKU1 and RV specific RNA detected.
+	-	-	+/-*	Only NL63/229E specific RNA detected.
-	+	-	+/-*	Only OC43/HKU1 specific RNA detected.
-	-	+**	+/-*	Only RV specific RNA detected.
-	-	-	+	Neither NL63/229E, nor OC43/HKU1, nor RV specific RNA detected. The sample does not contain detectable amounts of NL63/229E, OC43/HKU1 or RV specific RNA.
-	-	-	-	RT-PCR inhibition or reagent failure. Repeat testing from original sample or collect and test a new sample.

\* Detection of the IC in the JOE<sup>™</sup> detection channel is not required for positive results in the FAM<sup>™</sup> and/or the ROX<sup>™</sup> and/or the Cy5 detection channel. A high NL63/229E and/or OC43/HKU1 and/or RV RNA load in the sample can lead to reduced or absent IC signals.

\*\*Cross-reactivity of the rhinovirus specific detection system with some strains of enterovirus D and enterovirus C cannot be ruled out. These strains will lead to a weak signal in the rhinovirus detection channel (Cy5).

## 9. Technical assistance

For customer support, contact altona Diagnostics technical support:

e-mail:	support@altona-diagnostics.com
phone:	+49-(0)40-5480676-0

## 10. Trademarks and disclaimers

AltoStar<sup>®</sup>, FlexStar<sup>®</sup> (altona Diagnostics); QuantStudio<sup>™</sup> (Applied Biosystems); CFX96<sup>™</sup> (Bio-Rad); Rotor-Gene<sup>®</sup> (QIAGEN); FAM<sup>™</sup>, JOE<sup>™</sup>, ROX<sup>™</sup> (Thermo Fisher Scientific).

Registered names, trademarks, etc. used in this document, even if not specifically marked as such, are not to be considered unprotected by law.

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## 11. Explanation of symbols

Symbol	Explanation		
RUO	Research use only		
LOT	Batch code		
CONT	Content		
CAP	Cap color		
REF	Catalogue number		
NUM	Number		
СОМР	Component		
<b>[</b> ]i	Consult instructions for use		
Σ	Contains sufficient for "n" tests/reactions (rxns)		
X	Temperature limit		
$\mathbf{\Sigma}$	Use-by date		
	Manufacturer		
ΜΑΤ	Material number		

Symbol	Explanation
	Version
i	Note: Information is given to the user that is useful but not essential to the task at hand.

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