

## Instructions for use

# FlexStar<sup>®</sup> HAdV & HAstV & SaV RT-PCR Detection Mix 1.5

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Gastrointestinal

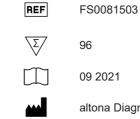


# **FlexStar**<sup>®</sup>

# HAdV & HAstV & SaV RT-PCR Detection Mix 1.5

For research use only!

(RUO)



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## 1. Application

The FlexStar<sup>®</sup> HAdV & HAstV & SaV RT-PCR Detection Mix 1.5 is a reagent system, based on real-time PCR technology, for the qualitative detection and differentiation of human adenovirus specific DNA, human astrovirus specific RNA and sapovirus specific RNA.

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

## 2. Product content

The FlexStar<sup>®</sup> HAdV & HAstV & SaV RT-PCR Detection Mix 1.5 contains the following components:

#### Table 1: Kit components

Lid color	Component	Number of tubes	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

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

<sup>2)</sup> Positive Control (HAdV specific DNA, HAstV specific RNA and SaV specific RNA)

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

## 3. Storage

- The FlexStar<sup>®</sup> HAdV & HAstV & SaV 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® HAdV & HAstV & SaV RT-PCR Detection Mix 1.5 is a reagent system. Used in combination with the FlexStar® (RT-)PCR Amplification Mix 1.5 it allows the qualitative detection and differentiation of human adenovirus (HAdV) specific DNA, human astrovirus (HAstV) specific RNA and sapovirus (SaV) specific RNA.

The FlexStar® HAdV & HAstV & SaV RT-PCR Detection Mix 1.5 is based on realtime (RT-)PCR technology, utilizing reverse-transcriptase (RT) reaction to convert HAstV and SaV RNA into complementary DNA (cDNA), polymerase chain reaction (PCR) for the amplification of HAdV, HAstV and SaV specific target sequences and fluorescently labeled target specific probes for the detection of the amplified DNA and cDNA, respectively. In addition to the HAdV specific DNA and the HAstV and SaV specific RNA specific amplification and detection systems the FlexStar® HAdV & HAstV & SaV RT-PCR Detection Mix 1.5 includes oligonucleotides for the amplification and detection of the internal control (IC, AltoStar® Internal Control 1.5). The IC is automatically added at the beginning of the nucleic acid purification procedure on the AltoStar® Automation System AM16 (in the following summarized as AltoStar® AM16). For details refer to the instructions for use of the AltoStar® Internal Control 1.5.

Probes specific for HAdV DNA are labeled with the fluorophore Cy5, probes specific for HAstV RNA are labeled with the fluorophore FAM<sup>™</sup> and probes specific for SaV RNA are labeled with the fluorophore ROX<sup>™</sup>, 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 HAdV, HAstV, SaV and the IC in the corresponding detection channels of the real-time PCR instrument.

## 4.1 Components

The FlexStar<sup>®</sup> HAdV & HAstV & SaV 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 (HAdV specific DNA, HAstV specific RNA and SaV 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 HAdV specific DNA, HAstV and SaV specific RNA, as well as of IC specific nucleic acids.

The PC contains HAdV specific DNA and HAstV as well as SaV specific RNA. It is used to verify the functionality of the HAdV, HAstV and SaV specific amplification and detection systems.

The NTC contains neither HAdV specific DNA, nor HAstV or SaV specific RNA but does contain the IC template. The NTC is used as negative control for the HAdV, the HAstV and the SaV specific real-time PCR and indicates possible contamination of the Detection Mix component.

## 4.2 Real-time PCR instruments

The FlexStar<sup>®</sup> HAdV & HAstV & SaV RT-PCR Detection Mix 1.5 can be used with the following real-time PCR instruments:

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

#### NOTE

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> HAdV & HAstV & SaV 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> HAdV & HAstV & SaV RT-PCR Detection Mix 1.5:

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

## 6. Procedure

#### 6.1 Sample preparation

Extracted nucleic acids are the starting material for the FlexStar<sup>®</sup> HAdV & HAstV & SaV RT-PCR Detection Mix 1.5. The quality of the extracted nucleic acids 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> HAdV & HAstV & SaV 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

#### 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		
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> HAdV & HAstV & SaV 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	

### 7.2 Fluorescence detectors (dyes)

Define the fluorescence detectors (dyes):

#### Table 5: Fluorescence detectors

Target	Detector name	Reporter	Quencher
HAdV specific DNA	HAdV	Cy5	(None)
HAstV specific RNA	HAstV	FAM™	(None)
SaV specific RNA	SaV	ROX™	(None)
Internal Control	IC	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:sec]
Reverse transcription	Hold	1	-	52	05:00
Denaturation	Hold	1	-	95	00:05
Amplification	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> HAdV & HAstV & SaV RT-PCR Detection Mix 1.5 on different real-time 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				
Cy5 (HAdV)	FAM™ (HAstV)	ROX™ (SaV)	JOE™ (IC)	Result interpretation
+	+	-	+/-*	HAdV specific DNA and HAstV specific RNA detected.
+	-	-	+/-*	Only HAdV specific DNA detected.
-	+	-	+/-*	Only HAstV specific RNA detected.
-	-	+	+/-*	Only SaV specific RNA detected.
+	-	+	+/-*	HAdV specific DNA and SaV specific RNA detected.
-	+	+	+/-*	HAstV specific RNA and SaV specific RNA detected.
+	+	+	+/-*	HAdV specific DNA, HAstV specific RNA and SaV specific RNA detected.
-	-	-	+	Neither HAdV specific DNA, nor HAstV specific RNA, nor SaV specific RNA detected. The sample does not contain detectable amounts of HAdV specific DNA, HAstV specific RNA or SaV 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™ detection channel is not required for positive results in the Cy5 and/or the FAM™ and/or the ROX™ detection channel. A high HAdV DNA and/or HAstV RNA and/or SaV RNA load in the sample can lead to reduced or absent IC signals.

## 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); JOE<sup>™</sup> (Life Technologies); Rotor-Gene<sup>®</sup> (QIAGEN); FAM<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
COMP	Component
Ĩ	Consult instructions for use
Σ	Contains sufficient for "n" tests/reactions (rxns)
X	Temperature limit
$\Sigma$	Use-by date
	Manufacturer
MAT	Material number
$\square$	Version
i	Note: Information is given to the user that is useful but not essential to the task at hand.
BIO	Contains biological material of animal origin

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