

Instructions for use

AltoStar[®] HAV RT-PCR Kit 1.5

12/2025 EN

AltoStar[®]

HAV RT-PCR Kit 1.5

For use with

ABI Prism[®] 7500 SDS (Applied Biosystems)

CFX96[™] Deep Well Dx System (Bio-Rad)

CFX96[™] Dx System (Bio-Rad)

LightCycler[®] 480 Instrument II (Roche)

QuantStudio[™] 5 Real-Time PCR System (Applied Biosystems)

Rotor-Gene[®] Q5/6 plex Platform (QIAGEN)



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1. About these instructions for use

These instructions for use guide the user in utilizing the AltoStar® HAV RT-PCR Kit 1.5. Chapters 1–6 and 9–14 contain general information and instructions which apply to each workflow used with the AltoStar® HAV RT-PCR Kit 1.5. Chapter 7 gives instructions on how to use the product with the AltoStar® Automation System AM16 (Hamilton; in the following summarized as AltoStar® AM16) with the AltoStar® Connect software (Version 1.7.4 or higher, Hamilton) for automated PCR setup and on the CFX96™ Deep Well Dx System (Bio-Rad, in the following summarized as CFX96™ DW Dx) with the CFX Manager™ Dx software (Version 3.1, Bio-Rad) for real-time PCR. Chapter 8 gives instructions regarding how to use the AltoStar® HAV RT-PCR Kit 1.5 in conjunction with other real-time PCR instruments. For details on the use of the AltoStar® AM16, the AltoStar® Connect software, the AltoStar® Purification Kit 1.5, the AltoStar® Internal Control 1.5 and the CFX96™ DW Dx, refer to the respective instructions for use listed below:

- AltoStar® Automation System AM16 Operator's Manual IVD (Hamilton)
- AltoStar® Connect Software Manual IVD (Hamilton)
- Instructions for use AltoStar® Purification Kit 1.5
- Instructions for use AltoStar® Internal Control 1.5
- CFX96™ Dx and CFX96™ Deep Well Dx Systems Operation Manual (Bio-Rad)

Throughout this manual, the terms CAUTION and NOTE have the following meanings:

CAUTION



Highlights operating instructions or procedures which, if not followed correctly, may result in personal injury or impact product performance. Contact altona Diagnostics technical support for assistance.

NOTE



Information is given to the user that is useful but not essential to the task at hand.

Read the instructions for use carefully before using the product.

2. Intended use

The AltoStar® HAV RT-PCR Kit 1.5 is an *in vitro* diagnostic test, based on real-time PCR technology, for the qualitative detection of hepatitis A virus (HAV) specific RNA (genotypes I – III) in human EDTA and citrate plasma. It is intended to be used as an aid for diagnosis of HAV infection.

The results generated with the AltoStar® HAV RT-PCR Kit 1.5 have to be interpreted in conjunction with other clinical and laboratory findings.

The AltoStar® HAV RT-PCR Kit 1.5 is intended for use by professional users trained in molecular biological techniques and *in vitro* diagnostic procedures.

3. Kit components

The AltoStar® HAV RT-PCR Kit 1.5 consists of a box with reagents and the AltoStar® HAV RT-PCR Kit 1.5 assay plugin.

3.1 Reagents

The AltoStar® HAV RT-PCR Kit 1.5 box contains the following components:

Table 1: Kit components

Lid color	Component	Number of tubes	Nominal volume [µl/tube]
Blue	Master A ¹⁾	8	60 ²⁾
Purple	Master B ¹⁾	8	180 ³⁾
Red	PC ⁴⁾	2	250
White	NTC ⁵⁾	2	250

¹⁾ Contains biological material of animal origin

²⁾ Contains an additional volume of 25 µl to compensate the dead volume of the AltoStar® AM16's liquid handling

³⁾ Contains an additional volume of 55 µl to compensate the dead volume of the AltoStar® AM16's liquid handling

⁴⁾ Positive Control

⁵⁾ No Template Control (negative control)

CAUTION



Before first use check the product and its components for completeness with respect to number, type and filling. Do not use a defective or incomplete product, product performance could be compromised.

The AltoStar® HAV RT-PCR Kit 1.5 contains enough reagents to perform 96 reactions in a maximum number of 8 runs.

The kit box is shipped on dry ice. Upon receipt and before first use, check the kit box and its components for:

- Integrity
- Completeness with respect to number, type and filling
- Correct labeling
- Expiration date
- Frozen state
- Clarity and absence of particles

If one or more product components are not frozen upon receipt, if tubes have been compromised during shipment or are missing, contact altona Diagnostics technical support for assistance (see chapter 12. Technical support).

3.2 AltoStar® HAV RT-PCR Kit 1.5 assay plugin

The AltoStar® HAV RT-PCR Kit 1.5 assay plugin version 1.0 can be optionally used in combination with the FastFinder Standalone software for data analysis and automated result interpretation. It is not provided with the AltoStar® HAV RT-PCR Kit 1.5 reagents and must be ordered separately as part of the AltoStar® Automated Analysis Package (Order No. VK000045).

4. Storage and handling

All reagents included in the AltoStar® HAV RT-PCR Kit 1.5 are ready-to-use solutions.

4.1 Storage

All reagents of the AltoStar® HAV RT-PCR Kit 1.5 must be stored at -25 °C to -15 °C upon arrival.

CAUTION



Improper storage conditions could compromise product performance.

CAUTION



Do not use products beyond the expiration date. The use of expired products could compromise product performance.

4.2 Handling

CAUTION



Do not exceed thaw-freeze-sequence and handling durations specified in these instructions for use, as this could compromise product performance.

CAUTION



Improper handling of product components and samples may cause contamination and could compromise product performance:

- Do not interchange vial or bottle caps.
- Store positive and/or potentially positive material separated from the kit components.
- Use separated working areas for sample preparation/reaction setup and amplification/detection activities.
- Always dispose gloves after handling positive and/or potentially positive material.
- Do not open the PCR plates and/or tubes post amplification.

CAUTION



Do not mix components from different kit lots. The use of different kit lots could compromise product performance.

4.2.1 Master A and Master B

After thawing, Master A and Master B are stable for 5 hours at up to +30 °C.

NOTE



If Master A and Master B were thawed but not used, they can be refrozen and thawed again once for later runs. If opened, discard the lids and use new lids to avoid contamination of the reagents.

4.2.2 Positive Control and No Template Control

1. After thawing, the Positive Control (PC) and the No Template Control (NTC) are stable for 5 hours at up to +30 °C.
2. Discard the lids of the PC and NTC tubes at each use and use new lids to avoid contamination of the reagents.
3. After use close the PC and NTC tubes with new lids and freeze them immediately.
4. Do not exceed the following thaw-freeze-sequence for each PC and NTC tube:
Thaw 1 → Freeze 1 → Thaw 2 → Freeze 2 → Thaw 3 → Freeze 3 → Thaw 4

5. Product description

The AltoStar® HAV RT-PCR Kit 1.5 is an *in vitro* diagnostic test for the qualitative detection of HAV specific RNA in human EDTA and citrate plasma.

It 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 HAV specific target sequences and fluorescently labeled target specific probes for the detection of the amplified DNA.

In addition to the HAV RNA specific amplification and detection systems, which target the gs1, gp1, gp2 (3C/3D junction region of the polyprotein) region of the viral genome, the assay includes oligonucleotides for the amplification and detection of the internal control (in the following abbreviated as IC; AltoStar® Internal Control 1.5).

The probe specific for HAV RNA is labeled with the fluorophore FAM™. The probe specific for the IC is labeled with a fluorophore (JOE™) detectable in the e.g., VIC™ channel.

Using probes linked to distinguishable dyes enables the parallel detection of HAV specific RNA and the IC in corresponding detection channels of the real-time PCR instrument used.

5.1 Background information

Hepatitis A virus (HAV) is one of five hepatitis viruses known to infect humans. It belongs to the family of *Picornaviridae* and the genus *Hepatovirus*. It is a non-enveloped and icosahedral virus and its genome consists of linear, positive-sense single-stranded RNA (ssRNA). HAV total genome size is approximately 7.5 kb in length.

HAV is primarily transmitted by the fecal-oral route through contact or smear infection. It is usually spread by drinking contaminated water or eating contaminated food (like shellfish), as well as by close contact with an already infected person. The virus is distributed worldwide. In developing countries nearly all individuals have a high risk of infection and had contact to HAV during their childhood. In more industrial developed regions like Europe and North America, infection risk is much lower since better sanitary standards are established. Following HAV infection permanent immunity is obtained.

After infection, first symptoms show after 14–28 days. Typical symptoms are fever, malaise, loss of appetite, diarrhea, nausea, dark colored urine and jaundice. Children that are infected often show no or only mild symptoms. However, adults often experience (severe) symptoms. In general mortality is low and infections with HAV does not turn into a chronic disease of the liver. There is no therapeutic treatment known for HAV infection but preventive vaccination is possible [1,2].

5.2 Reagent description

5.2.1 Master A and Master B

Master A and Master B contain all components (PCR buffer, reverse transcriptase, DNA polymerase, magnesium salt, primers and probes) to allow reverse transcription, as well as the PCR mediated amplification and target detection of HAV specific RNA and of the IC in one reaction setup.

5.2.2 Positive Control

The PC contains standardized concentrations of HAV specific RNA. It is calibrated against the 3rd WHO International Standard for Hepatitis A Virus VL For Nucleic Acid Amplification Techniques (NIBSC code: 15/276). The PC is used to verify the functionality of the HAV specific amplification and detection systems included in the AltoStar® HAV RT-PCR Kit 1.5.

5.2.3 No Template Control

The NTC contains no HAV specific RNA but does contain the IC template. The NTC is used as negative control for the HAV RNA specific real-time RT-PCR and indicates possible contamination of Master A and Master B.

5.3 Workflows

5.3.1 AltoStar® Workflow

The AltoStar® Workflow comprises the following products:

- AltoStar® Automation System AM16 (Hamilton)
- AltoStar® Connect software version 1.7.4 or higher (Hamilton)
- CFX96™ Deep Well Dx System (Bio-Rad) with CFX Manager™ Dx software version 3.1 (Bio-Rad)
- *Optional:* FastFinder Standalone version 1.0 or higher (UgenTec NV) and the AltoStar® HAV RT-PCR Kit 1.5 assay plugin for data analysis and automated result interpretation. The valid assay plugin version is specified in chapter 3.2 AltoStar® HAV RT-PCR Kit 1.5 assay plugin.

The workflow includes the following steps:

1. Programming an AltoStar® run.
2. Purification run on the AltoStar® AM16 using the AltoStar® Purification Kit 1.5 and the AltoStar® Internal Control 1.5.
3. RT-PCR setup run on the AltoStar® AM16 using the AltoStar® HAV RT-PCR Kit 1.5.
4. Real-time RT-PCR run on a CFX96™ DW Dx.

For further details on steps 3 and 4 of the workflow, refer to chapter 7. Using the AltoStar® HAV RT-PCR Kit 1.5 with the AltoStar® Workflow. All sample types and sample volumes specified for use with the AltoStar® Purification Kit 1.5 can be processed simultaneously on the AltoStar® AM16. Each sample can be analyzed with as many real-time PCR assays in parallel as the available eluate allows.

NOTE



Assays with differing PCR temperature profiles are automatically sorted to separate PCR plates.

5.3.2 Other workflows

The AltoStar® HAV RT-PCR Kit 1.5 can be used with compatible workflows. The following real-time PCR instruments and software have been validated for use with the AltoStar® HAV RT-PCR Kit 1.5:

- ABI Prism® 7500 SDS and the 7500 Software v2.3 (Applied Biosystems)
- CFX96™ Deep Well Dx System and the CFX Manager™ Dx software version 3.1 (Bio-Rad)
- CFX96™ Dx System and the CFX Manager™ Dx software version 3.1 (Bio-Rad)
- LightCycler® 480 Instrument II and the LightCycler® 480 Software version 1.5.1 (Roche)
- QuantStudio™ 5 Real-Time PCR System and the QuantStudio™ 5 Dx Software v1.0.2 (Applied Biosystems)

- Rotor-Gene® Q5/6 plex Platform and the Rotor-Gene® Q Software version 2.3.1 (QIAGEN)

Using one of the real-time PCR instruments listed above (except for the CFX96™ Deep Well Dx System), the PCR setup, the programming of the instrument as well as the data analysis have to be performed manually (for details see chapters 8.2.2 Master mix setup to 8.2.5 Data analysis).

5.4 Samples

5.4.1 Sample types

The following sample types are validated for use with the AltoStar® HAV RT-PCR Kit 1.5:

- Human EDTA plasma
- Human citrate plasma

CAUTION



This test has only been validated for use with human EDTA and citrate plasma. The use of other sample types could compromise product performance.

5.4.2 Sample collection and handling

Blood has to be collected with commercially available standard blood collection systems (e.g., Sarstedt, Becton Dickinson, Greiner or equivalent). Tube contents should be mixed directly after sample collection. The blood samples should be shipped cooled at +2 °C to +8 °C. Transport should occur following the local and national instructions for the transport of biological material.

For generation of EDTA or citrate plasma, whole blood should be centrifuged according to the instructions provided by the manufacturer of the collection system within 24 hours after collection. Before use EDTA or citrate plasma should not be stored for more than 48 hours at room temperature (+20 °C to +25 °C), 5 days at +2 °C to +8 °C or 2 months at -25 °C to -15 °C.

CAUTION



Always treat samples as infectious and (bio-)hazardous material in accordance with safety and laboratory procedures. For sample material spills promptly use an appropriate disinfectant. Handle contaminated materials as biohazardous.

CAUTION



Improper sample collection, shipment, storage or preparation could compromise product performance.

NOTE



Frozen storage of samples does not compromise kit performance. When working with frozen samples, make sure samples are completely thawed and properly mixed before use.

6. Warnings, precautions and limitations

- Before first use check the product and its components for completeness with respect to number, type and filling. Do not use a defective or incomplete product, product performance could be compromised.
- Improper storage conditions could compromise product performance.
- Do not use products beyond the expiration date. The use of expired products could compromise product performance.
- Do not exceed thaw-freeze-sequence and handling durations specified in these instructions for use, as this could compromise product performance.
- Improper handling of product components and samples may cause contamination and could compromise product performance:
 - Do not interchange vial or bottle caps.
 - Store positive and/or potentially positive material separated from the kit components.
 - Use separated working areas for sample preparation/reaction setup and amplification/detection activities.
 - Always dispose gloves after handling positive and/or potentially positive material.

- Do not open the PCR plates and/or tubes post amplification.
- Do not mix components from different kit lots. The use of different kit lots could compromise product performance.
- This test has only been validated for use with human EDTA and citrate plasma. The use of other sample types could compromise product performance.
- Always treat samples as infectious and (bio-)hazardous material in accordance with safety and laboratory procedures. For sample material spills promptly use an appropriate disinfectant. Handle contaminated materials as biohazardous.
- Improper sample collection, shipment, storage or preparation could compromise product performance.
- Storage of eluates under wrong conditions may lead to degradation of the HAV target sequence and could compromise product performance.
- Do not use an assay protocol version other than the one indicated on the 2D barcode in these instructions for use. Using an incorrect assay protocol version could compromise product performance.
- A lack of centrifugation of the product components after thawing may cause contamination with reagent residues in the lids and could compromise product performance.
- Do not reuse tube caps to avoid contamination of the reagents, as this could compromise product performance.
- As with any diagnostic test, results shall be interpreted in consideration of all clinical and laboratory findings.
- The presence of PCR inhibitors (e.g., heparin) could cause false negative or invalid results.
- Do not use different volumes of Master A and Master B for master mix setup than specified in these instructions for use, as this could compromise product performance.
- Do not exceed the PCR mix storage time, as this could compromise product performance.
- Do not mix up samples or sample IDs during PCR setup or transfer to the PCR instrument. This could lead to false positive or false negative results due to incorrect assignment of the samples.
- Do not use other cycling conditions than specified in these instructions for use, as this could compromise product performance.

- Do not use other control settings for data analysis than specified in these instructions for use, as this could cause incorrect IVD examination results.
- In case the sample contains other pathogens than HAV, competition with the target amplification or cross-reactivities may occur, causing incorrect IVD examination results.
- Disposal of hazardous and biological waste shall comply with local and national regulations to avoid environmental contamination.
- Potential mutations within the target regions of the HAV genome covered by primers and/or probes used in the kit may result in failure to detect the presence of the pathogen.
- This product is not intended to be used to detect the presence of HAV in blood, blood components, cells, tissues, organs or any of their derivatives in order to assess their suitability for transfusion, transplantation or cell administration.

7. Using the AltoStar® HAV RT-PCR Kit 1.5 with the AltoStar® Workflow

The following part of these instructions for use describes the use of the AltoStar® HAV RT-PCR Kit 1.5 in conjunction with the AltoStar® Workflow. The AltoStar® Workflow comprises different products [AltoStar® AM16, the AltoStar® Connect software, the AltoStar® Purification Kit 1.5, the AltoStar® Internal Control 1.5, the CFX96™ DW Dx, FastFinder Standalone and the AltoStar® HAV RT-PCR Kit 1.5 assay plugin for data analysis and automated result interpretation (optional)]. Use of those products is described in detail in the respective instructions for use.

- AltoStar® Automation System AM16 Operator's Manual IVD (Hamilton)
- AltoStar® Connect Software Manual IVD (Hamilton)
- Instructions for use AltoStar® Purification Kit 1.5
- Instructions for use AltoStar® Internal Control 1.5
- CFX96™ Dx and CFX96™ Deep Well Dx Systems Operation Manual (Bio-Rad)
- Instructions for use FastFinder Standalone (UgenTec NV)

7.1 Sample volume

The AltoStar® HAV RT-PCR Kit 1.5 is validated for nucleic acid purifications from a sample volume of 500 µl when using the AltoStar® AM16. Additional sample volume has to be provided to account for the dead volume of the sample tube used (see chapter 7.2 Sample tubes).

7.2 Sample tubes

Sample tubes suitable for use on the AltoStar® AM16 can be purchased from altona Diagnostics (7 ml tube with cap, 82 x 13 mm, VK000010). Other sample tubes can be tested for applicability by the user. For details, refer to the instructions for use of the AltoStar® Purification Kit 1.5.

7.3 Sample barcodes

For automated sample identification by the AltoStar® AM16, all sample tubes must be labeled with a suitable barcode. For details, refer to the instructions for use of the AltoStar® Purification Kit 1.5.

7.4 Material and devices required but not provided for the AltoStar® Workflow

The material and devices shown in table 2 must be ordered from altona Diagnostics.

Table 2: Required material and devices

Material	Description	Order No.
AltoStar® Molecular Diagnostic Workflow	Product bundle containing the AltoStar® Automation System AM16, the AltoStar® Connect software (Version 1.7.4 or higher) and IT hardware	AM16
AltoStar® Detection	Product bundle containing the CFX96™ Deep Well Dx System with CFX Manager™ Dx software (Version 3.1), a barcode scanner and IT hardware	DT16

Material	Description	Order No.
AltoStar® Purification Kit 1.5	Nucleic acid isolation and purification chemistry for use with the AltoStar® Automation System AM16	PK15-46
AltoStar® Internal Control 1.5	Nucleic acid extraction and PCR amplification and detection control	IC15-46
AltoStar® Automated Analysis Package	Personal computer with installed FastFinder Standalone software (Version 1.0 or higher) and assay plugin bundle for data analysis and automated result interpretation	VK000045
PCR Plate	Semi-skirted, barcoded 96 multi-well plate with white wells	VK000005
PCR Plate Sealing Foil	Sealing foil for the PCR plate	VK000006
1,000 µl CO-RE Tips	1,000 µl filter tips for use with the AltoStar® Automation System AM16	VK000007
300 µl CO-RE Tips	300 µl filter tips for use with the AltoStar® Automation System AM16	VK000008
Pooling Tube	Barcoded tube for pooling of PCR reagents	VK000002
Waste Bag	Autoclavable sterile bag for use with the AltoStar® Automation System AM16	VK000009
Screw Cap - red	Screw cap for PC tubes	VK000012
Screw Cap - blue	Screw cap for Master A tubes	VK000013
Screw Cap - purple	Screw cap for Master B tubes	VK000015
Screw Cap - white	Screw cap for NTC tubes	VK000016

Table 3: Additional laboratory material and devices

Material	Description	Order No.
Plate Sealer	e.g., AltoStar® Plate Sealer	VK000023
	e.g., PX1 PCR Plate Sealer (Bio-Rad)	VK000033

7.5 General material and devices

- Vortex mixer
- Powder-free gloves (disposable)
- Centrifuge for centrifugation of the AltoStar® HAV RT-PCR Kit 1.5 components
- Centrifuge for centrifugation of PCR plates

7.6 Procedure

7.6.1 Overview of the AltoStar® Workflow

The steps of the complete AltoStar® Workflow are summarized in table 4. Information on specific settings for use with the AltoStar® HAV RT-PCR Kit 1.5 are provided in chapter 7.6.2 Programming an AltoStar® run. For detailed instructions for steps 1–5, refer to the instructions for use of the AltoStar® Purification Kit 1.5, the AltoStar® Connect software and the AltoStar® AM16.

Steps 6–9 are described in more detail in chapters 7.6.3 Starting a PCR setup run to 7.6.6 Starting a PCR run.

Steps 10 and 11 differ for manual and automated result interpretation. More details on manual result interpretation can be found in chapters 7.6.7 PCR data analysis using CFX Manager™ Dx software to 7.6.10 Export of PCR results for manual result interpretation. The steps for optional data analysis and automated result interpretation using FastFinder are described in more detail in chapter 7.6.11 PCR data analysis and automated result interpretation using FastFinder.

Table 4: Overview of the AltoStar® Workflow

Step	Action
1. Start the AltoStar® AM16	<ul style="list-style-type: none"> • Switch on the AltoStar® AM16. • Switch on the computer and the monitor. • Start the AltoStar® Connect software.
2. Perform maintenance	<ul style="list-style-type: none"> • In the menu bar click Application → Instrument Maintenance. <ul style="list-style-type: none"> ◦ If weekly maintenance is due, click Start Weekly Maintenance. ◦ If daily maintenance is due, click Start Daily Maintenance. • Follow the on-screen instructions for the maintenance process.
3. Program an AltoStar® run	<ul style="list-style-type: none"> • In the menu bar click Program Run → Program Run (AltoStar® Purification). Alternatively, go back to the Start screen and click the Program Run button. • Enter samples or import from the LIMS. • Select the following assay for the samples unless already imported from the LIMS: <ul style="list-style-type: none"> ◦ AltoStar® HAV RT-PCR Kit 1.5 • Click the Create Run button in the tool bar to create the AltoStar® run.

Step	Action
<p>4. Start a purification run</p>	<ul style="list-style-type: none"> • In the menu bar click Purification → Start Purification. Alternatively, go back to the Start screen and click the Start Purification button. • Select the purification run to be started to display the samples included in the selected purification run. • Prepare the purification reagents: <ul style="list-style-type: none"> ◦ Ensure that the purification reagents to be used have the same loading number (except AltoStar® Internal Control 1.5) and are not expired. ◦ If precipitates are visible in the Lysis Buffer, heat it ($\leq +50$ °C) until completely dissolved. ◦ Thaw the IC (AltoStar® Internal Control 1.5) and vortex for 5 seconds. ◦ Vortex the Magnetic Beads for 5 seconds without wetting the lid. • Prepare the samples for the purification run to be started as described in the instructions for use of the AltoStar® Purification Kit 1.5. • Click the Start Run button in the tool bar. • Follow the loading dialogs and load the instrument accordingly. • Confirm the Loading complete message with OK or wait 10 seconds. <p>The system will now perform the purification run automatically.</p>
<p>5. Finish the purification run</p>	<ul style="list-style-type: none"> • Make sure the loading tray is empty and confirm the Run finished dialog with OK. • Follow the instructions in the Maintenance dialog and confirm with OK. • Seal and store the components of the AltoStar® Purification Kit 1.5 that can be reused. <p>The eluates in the unsealed eluate plate are stable at room temperature (max. +30 °C) for a total of 4 hours.</p> <ul style="list-style-type: none"> • If the associated PCR setup run is not started right away, seal the eluate plate with an eluate plate sealing foil and store at +2 °C to +8 °C for up to 24 hours. • View the purification run results to confirm successful processing of each sample.

Step	Action
<p>6. Start a PCR setup run</p>	<ul style="list-style-type: none"> • In the menu bar, click PCR Setup → Start PCR Setup. Alternatively, go back to the Start screen and click the Start PCR Setup button. • Select the PCR setup run to be started in order to display the eluate plate and reagents included in the selected PCR setup run. • Prepare the PCR reagents: <ul style="list-style-type: none"> ◦ Ensure that masters and controls to be used are from the same kit lot and are not expired. ◦ Thaw the required amount of master and control tubes, vortex briefly and spin down in a centrifuge. • If the eluate plate is sealed, briefly centrifuge the plate and unseal carefully. • Click the Start Run button in the tool bar. • Follow the Loading dialog and load the instrument accordingly. • Confirm the Loading complete message with OK or wait 10 seconds. <p>The system will now perform the PCR setup run automatically.</p>
<p>7. Finish the PCR setup run</p>	<ul style="list-style-type: none"> • Make sure the loading tray is empty and confirm the Run finished dialog with OK. • Follow the instructions in the Maintenance dialog and confirm with OK. • Close and store the components of the AltoStar® HAV RT-PCR Kit 1.5 that can be reused. • View the PCR setup run results to confirm successful processing of each sample.
<p>8. Seal the PCR plate</p>	<ul style="list-style-type: none"> • Seal the PCR plate with a PCR plate sealing foil.

Step	Action	
<p>9. Start the PCR run</p>	<ul style="list-style-type: none"> • Switch on the CFX96™ DW Dx, the attached computer and the monitor. • Start the CFX Manager™ Dx software. • Open the CFX96™ DW Dx. • Centrifuge the PCR plate and insert it into the CFX96™ DW Dx. • Select File → Open → LIMS File... from the menu bar. • Scan the barcode of the PCR plate with the handheld barcode scanner. • Close the CFX96™ DW Dx. • Click the Start Run button to start the PCR run. Name and save the PCR run file. <p>The CFX96™ DW Dx will now perform the PCR run automatically.</p>	
Step	Action for data analysis and report generation with CFX Manager™ Dx software for subsequent manual result interpretation or result interpretation by LIMS	Action for optional data analysis and automated result interpretation using FastFinder
<p>10. Separate assays for individual analysis</p>	<ul style="list-style-type: none"> • Separate all assays in the PCR run into distinct well groups. 	<p>Not required.</p>
<p>11. Analyze the data and interpret the PCR run results</p>	<p>For each well group individually:</p> <ul style="list-style-type: none"> • Perform baseline correction in all wells for all detection channels used. • Exclude wells with irregular PCR signals. • Set the thresholds of all detection channels according to the controls. • Exclude wells containing invalid data. • Generate the LIMS result file for export of results to the LIMS. • Generate the result report for manual result interpretation. 	<ul style="list-style-type: none"> • Use FastFinder Standalone and the AltoStar® HAV RT-PCR Kit 1.5 assay plugin for data analysis and automated result interpretation.

CAUTION

Storage of eluates under wrong conditions may lead to degradation of the HAV target sequence and could compromise product performance.

7.6.2 Programming an AltoStar® run

For detailed information on starting an AltoStar® run, refer to the instructions for use of the AltoStar® Purification Kit 1.5, the AltoStar® Connect software and the AltoStar® AM16. Specific settings for use with the AltoStar® HAV RT-PCR Kit 1.5 are listed below:

- PC and NTC are selected.
- The required sample volume is 500 µl plus the dead volume for the respective sample tube (see chapters 7.1 Sample volume and 7.2 Sample tubes).
- The required eluate volume for the AltoStar® HAV RT-PCR Kit 1.5 is 10 µl.
- Make sure that the correct assay protocol version is used for the run. For information on the current protocol version see chapter 17. Assay protocol for the AltoStar® Connect software and information for LIMS integration. The respective assay protocol is encoded in the 2D barcode displayed there. For information on purification and assay protocol import into the AltoStar® Connect software, refer to the respective instructions for use.

CAUTION

Do not use an assay protocol version other than the one indicated on the 2D barcode in these instructions for use. Using an incorrect assay protocol version could compromise product performance.

7.6.3 Starting a PCR setup run

1. Select **PCR Setup** → **Start PCR Setup** in the menu bar. Alternatively, go back to the Start screen of the AltoStar® Connect software and select the **Start PCR Setup** button. The Start PCR Setup Run screen is displayed.

The pending PCR setup runs are displayed in the Programmed PCR Setup Runs table on the left side of the screen.

2. Select the PCR setup run to be started in the Programmed PCR Setup Runs table.
 - The samples included in the selected PCR setup run are displayed in the table on the top right side of the screen (Samples in selected PCR Setup Run).
 - The controls required for the selected PCR setup run are displayed in the table on the middle right side of the screen (Controls in selected PCR Setup Run).
 - The number of master tubes required for the selected PCR setup run is displayed in the table on the bottom right side of the screen (Required master tubes for the selected PCR Setup Run).

NOTE

The number of prioritized samples in a PCR setup run is displayed in the column **No. of prioritized Samples**. Conduct PCR setup runs with prioritized samples first to facilitate fastest processing of prioritized samples.

Before clicking the **Start Run** button in the tool bar, prepare the required reagents as described in chapter 7.6.3.1 Preparing reagents for a PCR setup run. If the eluate plate required for the selected PCR setup run has been sealed for storage prepare it as described in the instructions for use of the AltoStar® Purification Kit 1.5.

7.6.3.1 Preparing reagents for a PCR setup run

1. Thaw the required controls and the required number of master tubes completely at room temperature (max. +30 °C).
2. Mix the reagents by gentle vortexing.
3. Centrifuge the tubes briefly to remove drops from the lid.

CAUTION

A lack of centrifugation of the product components after thawing may cause contamination with reagent residues in the lids and could compromise product performance.

7.6.3.2 Loading the AltoStar® AM16 for a PCR setup run

For detailed information regarding the loading process, refer to the instructions for use of the AltoStar® AM16 and the AltoStar® Connect software.

1. Click the **Start Run** button in the tool bar of the Start PCR Setup Run screen to display the Loading dialog.

The Loading dialog consists of a visual representation of the AltoStar® AM16 deck at the top and a table specifying the carriers, the respective tracks on the AltoStar® AM16 deck for each carrier, the material to be loaded onto each carrier and comments with respect to the carrier loading.

NOTE

To visualize the position of an item on a carrier and the position of the carrier on the AltoStar® AM16 deck, select the respective row of the table in the Loading dialog.



The position of the item and its carrier is visualized:

- Highlighted in red in the visual representation of the instrument deck
- On the AltoStar® AM16 by flashing loading lights above the tracks where the selected carrier must be placed

2. Load the required material, the prepared eluate plate and the prepared reagents onto the suitable carriers.
 - Exchange only **completely empty** 1,000 µl tip racks for **completely full** 1,000 µl tip racks on the tip carrier.
 - Exchange only **completely empty** 300 µl tip racks for **completely full** 300 µl tip racks on the tip and plate carrier.

NOTE



Exchange of tip racks, which are not completely empty as well as handling of individual tips may interfere with the automatic tip management and cause run aborts.

- Place the required eluate plate with well A1 to the left of the black plate position.

- Place a PCR plate with well A1 to the left of the silver front plate position.
- Load a tube carrier 24 with one unused pooling tube for each assay in the PCR setup run.
- Gently push the tubes all the way to the bottom of the carrier and rotate the tubes until the tube barcodes are visible through the carrier window.
- Load the reagent tube carrier 32 with the assay components required for the PCR setup run.
- Gently push the tubes all the way to the bottom of the carrier and rotate the tubes until the tube barcodes are visible through the carrier window.

NOTE



The position of the individual tubes on the carriers is arbitrary.

NOTE



The volume of the loaded components is not checked by the system prior to processing. Insufficient component volume will prevent successful PCR setup for the affected assay.

NOTE



Starting a PCR setup run with lids still on the tubes may cause the run to abort during processing.

3. Load the carriers with the carrier barcode towards the rear facing right.
4. Insert populated carriers into the respective tracks between the front and rear slide blocks of the loading tray until they touch the stop hooks on the far side of the loading tray.

NOTE



Pushing the carriers past the stop hooks may damage the instrument and interfere with the loading process.

5. Check that the tip eject sheet and the tip waste container are in the correct position and a new waste bag is placed in the container.

- Click **OK** in the Loading dialog to proceed with the loading process.

NOTE

By clicking **Cancel** the PCR setup run will be canceled, but it can be started again (see chapter 7.6.3 Starting a PCR setup run).

The AltoStar® AM16 draws the carriers into the instrument and performs barcode verification.

NOTE

The AltoStar® AM16 automatically verifies:

- Correct type and localization of the loaded carriers
- Correct identity and position of the items loaded on the carriers
- Lot congruence of the components of the individual AltoStar® assay kits
- Non-expiry of all loaded AltoStar® assay components
- Correct positioning of the tip eject sheet

If any of these checks fail, the user is prompted with a message dialog specifying the problem at hand and instructions to correct the issue accordingly. For further information regarding error handling, refer to the instructions for use of the AltoStar® Connect software.

NOTE

Altering positions of any loaded item after the carrier has been drawn into the instrument may result in the abort of the PCR setup run and/or damage to the instrument.

When all checks have passed the Loading complete dialog is displayed.

- Confirm the Loading complete dialog by clicking **OK** or wait 10 seconds for the automatic start of the process.

NOTE

By clicking **Cancel** the PCR setup run will be canceled, but it can be started again (see chapter 7.6.3 Starting a PCR setup run).

The PCR setup run is started and will be conducted without user intervention.

7.6.3.3 During the PCR setup run

No further user interaction is required until the PCR setup run has finished. The Processing Status screen is displayed showing the status of the PCR setup run and the estimated time remaining.

NOTE

Pushing or pulling carriers or the door of the AltoStar® AM16 during a PCR setup run may abort the run.

7.6.4 End of the PCR setup run

At the end of the PCR setup run the Run finished dialog is displayed.

1. Make sure that the loading tray is empty.
2. Confirm the Run finished dialog by clicking **OK**.

The AltoStar® AM16 will unload the carriers. Make sure not to stand in the way of the unloading carriers.

After unloading the Maintenance dialog is displayed.

3. Follow the instructions of the Maintenance dialog.

The table of the dialog displays the number of reactions in the master tubes that were not used in the PCR setup run.

4. If another PCR setup run using the currently loaded eluate plate is to be started right away, the eluate plate can remain unsealed on the carrier position. If this is **not** the case, seal and store the eluate plate. For further information, refer to the instructions for use of the AltoStar® Purification Kit 1.5.

NOTE



The eluates in the eluate plate are stable at room temperature (max. +30 °C) for a total of 4 hours after completion of the purification run.

5. Close reagent tubes with suitable unused tube caps.

CAUTION



Do not reuse tube caps to avoid contamination of the reagents, as this could compromise product performance.

6. Store reagents for reuse as described in chapter 4.2 Handling.
7. Dispose of the used materials (see chapter 10. Disposal).
8. Confirm the Maintenance dialog by clicking **OK**.

7.6.4.1 PCR setup run results

The PCR setup run results are saved in the AltoStar® Connect software.

1. Click **PCR Setup** → **PCR Setup Results** in the menu bar to access the Results screen.

The Results screen displays a table with all samples used in the latest PCR setup run and a column **Status** at the right showing if the PCR setup process for a given sample was conducted completely (see table 5).

Table 5: PCR setup run results

Status	PCR setup run result
Processed	<ul style="list-style-type: none"> The eluate was successfully processed in the PCR setup run. The resulting RT-PCR mix is ready for use in a PCR run.
Error	<ul style="list-style-type: none"> The eluate was not processed successfully. The respective RT-PCR mix will be automatically omitted in the following PCR analysis.

- To view the results of prior PCR setup runs, click the **Load** button in the menu bar, select the desired PCR setup run from the list in the opening Load Results dialog and click **OK**.

3 PCR setup run result files are automatically generated by the AltoStar® Connect software:

- A LIMS file (.XML) to pass detailed information about the PCR setup run including results back to the LIMS
- A report (.PDF) containing detailed information about the PCR setup run including results for documentation purposes
- A cycler file (.PLRN) for automatic programming of the CFX96™ DW Dx

These files are saved to the location specified in the system settings of the AltoStar® Connect software.

NOTE



PCR setup run result files can be generated again by loading the respective PCR setup run and clicking the **Create LIMS File** button to generate the LIMS file, the **Create Report** button to generate the report or the **Create Bio-Rad Cycler File** button to generate the cycler file.

7.6.5 Sealing of the PCR plate

After completion of the PCR setup run, the PCR plate must be sealed with a PCR plate sealing foil. It is recommended to use the AltoStar® Plate Sealer [4s3™ Semi-Automatic Sheet Heat Sealer (4titude)] or the PX1 PCR Plate Sealer (Bio-Rad). The suitability of plate sealers other than the recommended plate sealers has to be evaluated by the user.

If one of the recommended plate sealers is used for sealing, proceed as follows:

1. Switch on the plate sealer and make sure that the plate adapter is not in the drawer.
2. Ensure that the settings of the plate sealer are as follows:

Table 6: Settings of the plate sealer

Plate sealer	Settings	
	Temperature [°C]	Time [s]
AltoStar® Plate Sealer [4s3™ Semi-Automatic Sheet Heat Sealer (4titude)]	170	2
PX1 PCR Plate Sealer (Bio-Rad)	175	3

3. Wait until the set temperature is reached. This may take several minutes.
4. Place the PCR plate on the plate adapter of the plate sealer.
5. Place one PCR plate sealing foil on the PCR plate so that the print 'THIS SIDE UP' is readable. Make sure that all wells of the PCR plate are covered with foil and no well is obscured by the writing.




NOTE



Operating the plate sealer without the plate adapter placed in the drawer may render the sealer nonfunctional. In this case, contact Altona Diagnostics technical support for assistance (see chapter 12. Technical support).

NOTE

If the PCR plate sealing foil or the frame is placed incorrectly, the foil may stick to the heating plate within the plate sealer during sealing. This will render the sealer nonfunctional. In this case, or if the sealing step has been initiated without a PCR plate sealing foil, let the plate sealer cool down to room temperature and contact Altona Diagnostics technical support for assistance (see chapter 12. Technical support).

6. Assemble the sealing frame on top to hold down the sealing foil.
7. Open the drawer by pressing the **Operate***/ ** button.
8. Place the assembly consisting of the plate adapter, the PCR plate, the PCR plate sealing foil and the sealing frame into the plate sealer and press the **Operate***/ ** button.
9. The drawer closes automatically, seals for the set time and reopens automatically.
10. Take the sealed PCR plate and the plate adapter out of the plate sealer and close the plate sealer by pressing the **Close***/ ** button.

* AltoStar® Plate Sealer [4s3™ Semi-Automatic Sheet Heat Sealer (4titude)]

**PX1 PCR Plate Sealer (Bio-Rad)

7.6.5.1 PCR mix stability

After completion of the PCR setup run the RT-PCR mix in the sealed PCR plate is stable at room temperature (max. +30 °C) for 30 minutes.

CAUTION

Do not exceed the PCR mix storage time, as this could compromise product performance.

7.6.6 Starting a PCR run

The PCR run is performed on a CFX96™ DW Dx under control of the CFX Manager™ Dx software.

1. Switch on the CFX96™ DW Dx, the attached computer and the monitor.

2. Start the CFX Manager™ Dx software.
3. In the menu bar of the CFX Manager™ Dx software select **File** → **Open** → **LIMS File...** to open the Open LIMS File dialog.
4. In the opening Open LIMS File dialog, make sure that the cursor is blinking in the **File name** field at the bottom. If it is not, click into the **File name** field.
5. Scan the PCR plate barcode with the handheld barcode scanner to automatically select and open the correct LIMS file. The Run Setup dialog is displayed.

NOTE

All parameters required for the start of the PCR run are transferred automatically from the AltoStar® Connect software to the CFX96™ DW Dx by means of the cyclers file.

6. Click the **Open Lid** button in the Run Setup dialog to open the lid of the CFX96™ DW Dx.
7. Briefly centrifuge the sealed PCR plate to ensure all liquid is at the bottom of the wells.
8. Insert the sealed PCR plate into the heating block of the CFX96™ DW Dx with well A1 to the left side.
9. Close the CFX96™ DW Dx by clicking the **Close Lid** button in the Run Setup dialog.
10. Start the PCR run by clicking the **Start Run** button in the Run Setup dialog.

7.6.6.1 During the PCR run

No user interaction is required until the PCR run has finished. The Run Details dialog is displayed showing the status of the PCR run and the estimated time remaining.

NOTE

Opening the lid of the CFX96™ DW Dx during a PCR run by operating the button at the front of the lid or by clicking **Open Lid** in the Run Details dialog will abort the run and void all results.

At the end of the PCR run the Data Analysis window is displayed, which shows the amplification curves, the plate layout and the results.

7.6.7 PCR data analysis using the CFX Manager™ Dx software

The results of all assays (well groups) on the PCR plate have to be analyzed in the sequence depicted in figure 1.

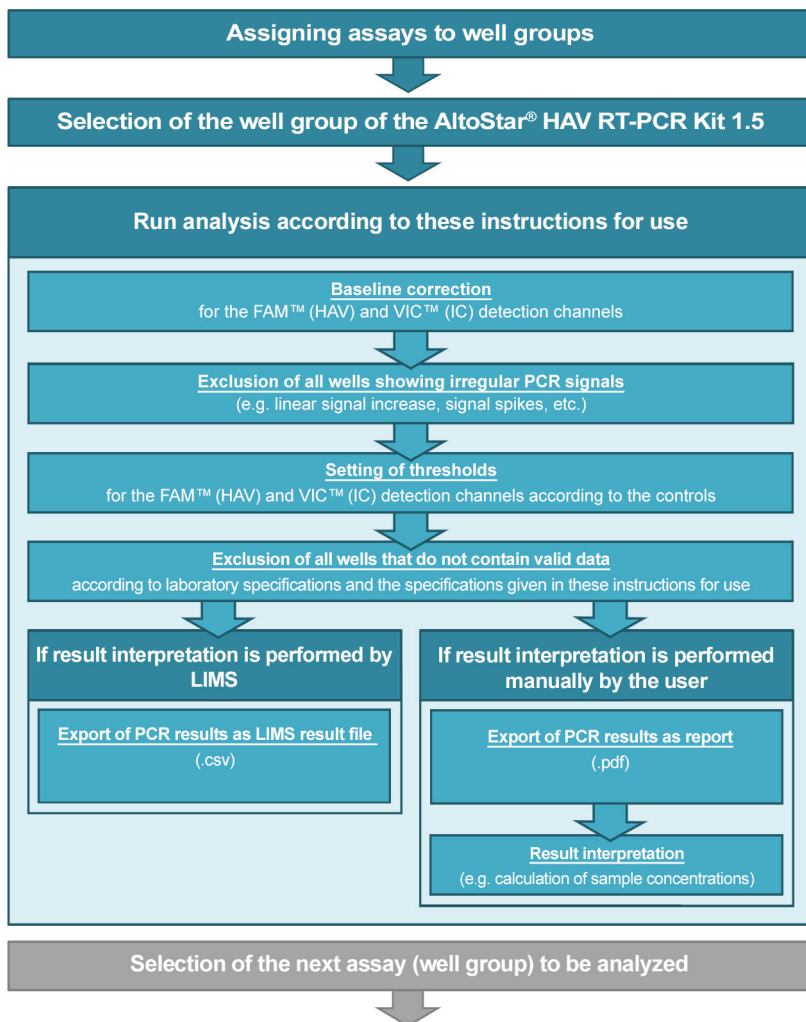


Figure 1: PCR data analysis process using the CFX Manager™ Dx software

7.6.7.1 Assigning assays to well groups

The AltoStar® Workflow processes one or several PCR assays simultaneously on one PCR plate. However, each assay must be analyzed separately by the user according to the instructions for use of the respective assay.

To this end, all assays on a PCR plate must be assigned to individual well groups in the CFX Manager™ Dx software by the user.

1. In the Data Analysis window click the **Plate Setup** button in the toolbar and select **View/Edit Plate**. The Plate Editor dialog is displayed (see figure 2).

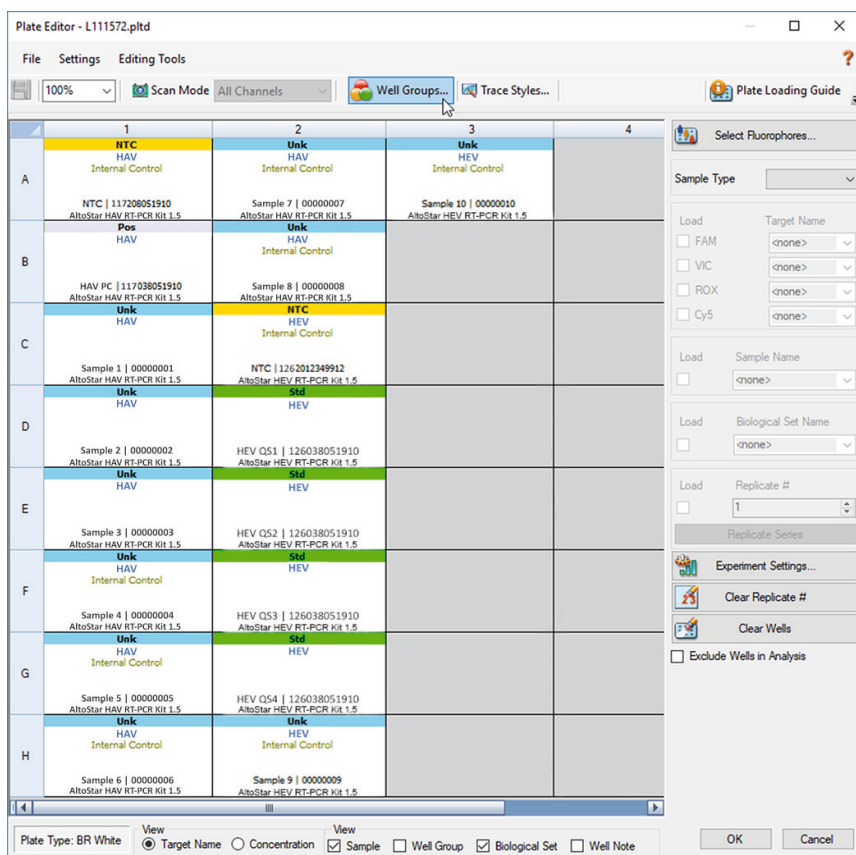


Figure 2: Plate Editor dialog

- In the Plate Editor dialog click **Well Groups...** in the toolbar. The Well Groups Manager dialog is displayed (see figure 3).
- Click the **Add** button.
- Type the name of the first assay in the text box.
- Select all wells in the PCR plate area that belong to the first assay (see figure 3). The wells belonging to an individual assay can be identified in the Plate Editor dialog by the entry in the **Biological Set** field.

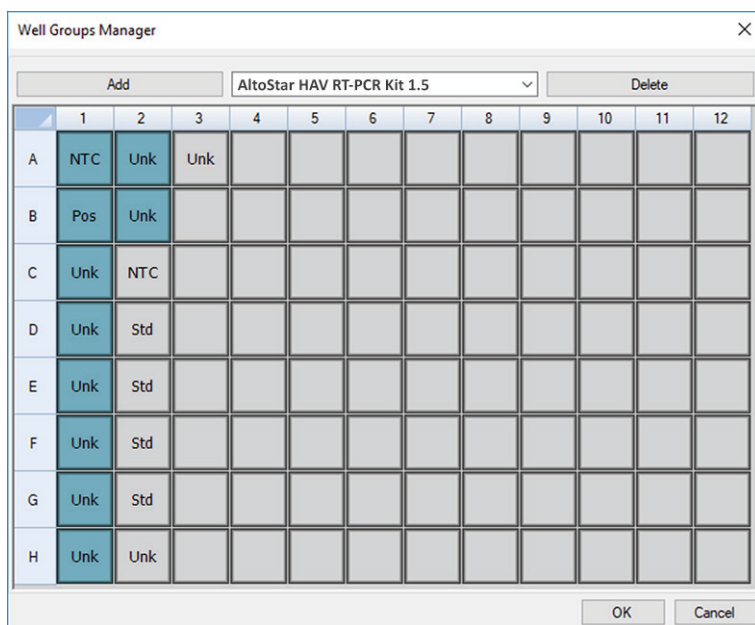


Figure 3: Well Groups Manager dialog

- Repeat steps 3–5 for all assays on the PCR plate.
- Confirm well group assignment by clicking **OK**. The Well Groups Manager dialog closes.
- Close the Plate Editor dialog by clicking **OK**.
- Confirm to apply the changes by clicking **Yes**.

In the Data Analysis window ensure to select the **Well Group** of the AltoStar® HAV RT-PCR Kit 1.5. Therefore, click on the **Well Group** drop-down menu right next to the **Well Group** button (see figure 4) of the toolbar. Do not use the “All Wells” **Well Group**. The selection in figure 4 is used as a general example view.

Before analyzing the results, ensure that the well group of the AltoStar® HAV RT-PCR Kit 1.5 contains all of the AltoStar® HAV RT-PCR Kit 1.5 wells and no wells of other assays.

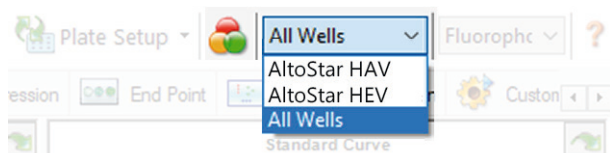


Figure 4: Well Group button and Well Group drop-down menu

NOTE



Combined analysis of more than one assay may lead to incorrect results.

CAUTION



As with any diagnostic test, results shall be interpreted in consideration of all clinical and laboratory findings.

7.6.7.2 Baseline correction

The baseline settings used by the CFX Manager™ Dx software may have to be corrected for individual wells of the assay (**Well Group**) under analysis.

1. In the Data Analysis window ensure to select the **Well Group** of the AltoStar® HAV RT-PCR Kit 1.5. Therefore, click on the **Well Group** drop-down menu right next to the **Well Group** button (see figure 4) of the toolbar.
2. At the left side of the Data Analysis window tick only the **FAM** checkbox for the HAV target detection channel.

- In the menu bar of the Data Analysis window click **Settings** → **Baseline Threshold...** to open the Baseline Threshold dialog (see figure 5).
- Click the \diamond symbol in the **Baseline End** column header once to sort the table by ascending **Baseline End** values.
- Select all lines that show a **Baseline End** value of 1–9 (see figure 5).

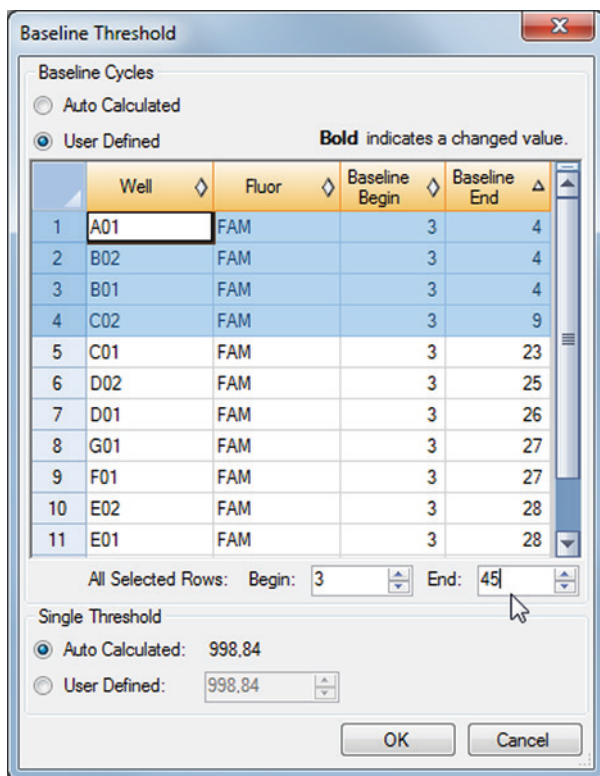


Figure 5: Baseline Threshold dialog

- Set the value in the **End:** field to 45 for the selected lines (see figure 5).
- Confirm the settings by clicking **OK**.
- At the left side of the Data Analysis window untick the **FAM** checkbox and tick only the **VIC** checkbox for the IC target detection channel.
- Repeat steps 3–7 for the VIC™ (IC) detection channel.

7.6.7.3 Exclusion of irregular PCR signals

Valid results can only be derived from PCR signals that are free of signal artefacts, which may be caused e.g., by impurities or bubbles in the RT-PCR mix. PCR signals that contain signal artefacts have to be excluded by the user.

1. In the Data Analysis window ensure to select the **Well Group** of the AltoStar® HAV RT-PCR Kit 1.5. Therefore, click on the **Well Group** drop-down menu right next to the **Well Group** button (see figure 4) of the toolbar.

- Identify wells with irregular PCR signals (linear signal increase, signal spikes, etc.) in any of the FAM™ (HAV target) and the VIC™ (IC) detection channels (see figure 6).

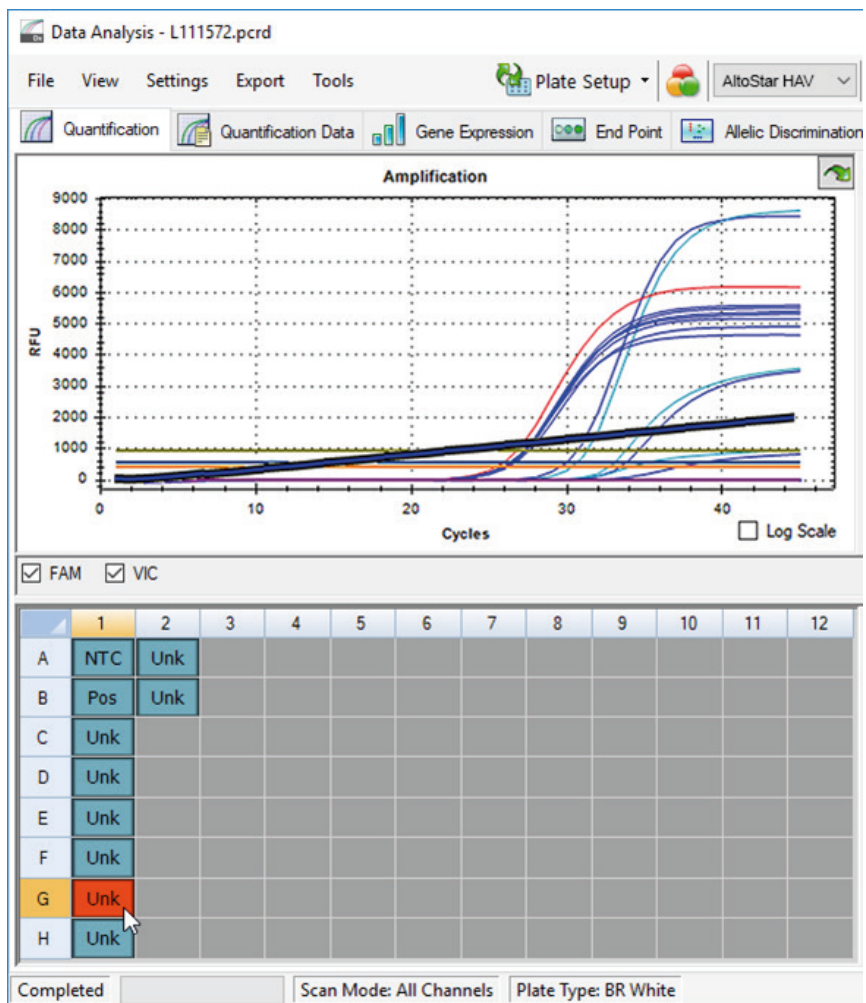


Figure 6: Data Analysis window: irregular PCR signal

- Click each affected well with the right mouse button and select **Well...** → **Exclude from Analysis** (see figure 7).

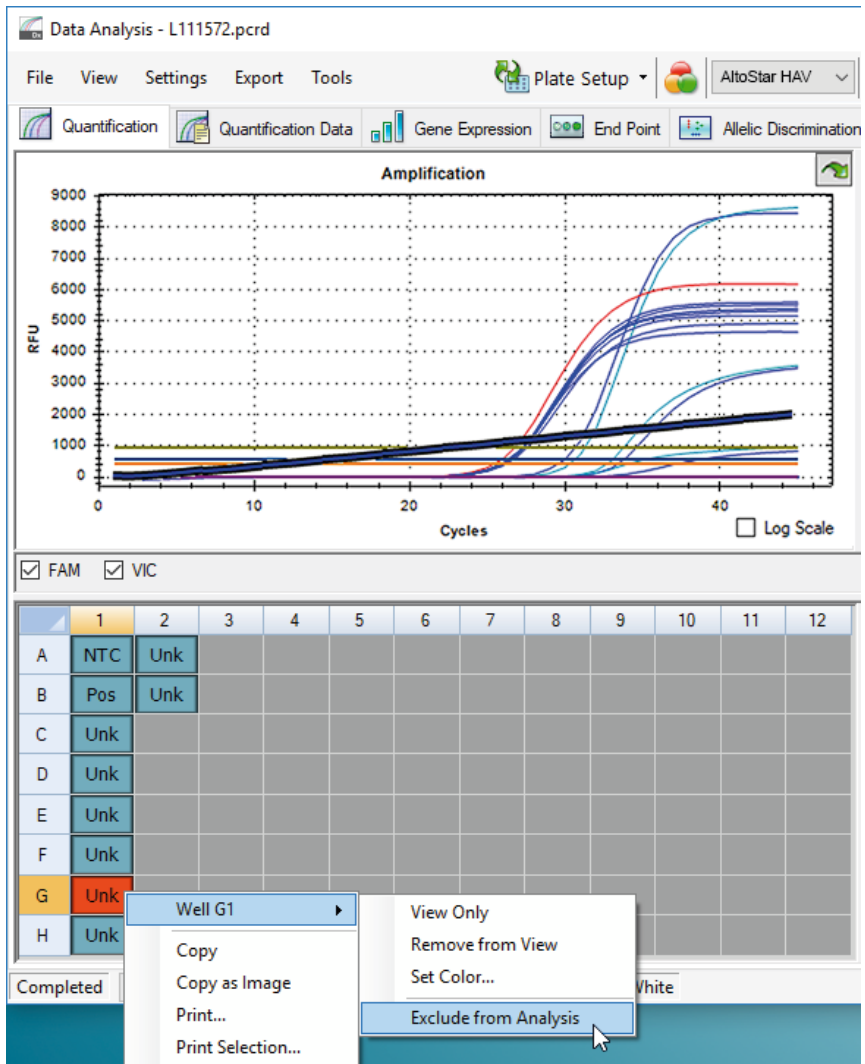


Figure 7: Data Analysis window: exclude well from analysis

4. The selected well is excluded from the analysis. No results will be generated for this well (see figure 8).

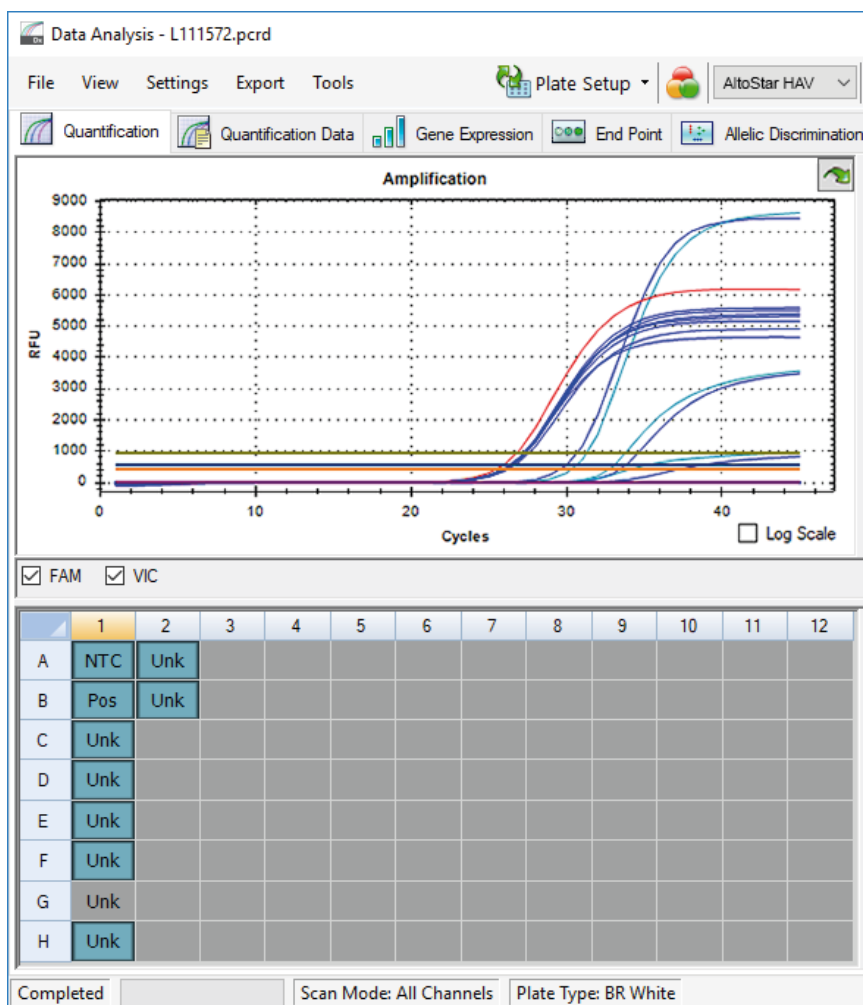


Figure 8: Data Analysis window: excluded well

7.6.7.4 Setting of thresholds

The thresholds for the FAM™ (HAV target) and the VIC™ (IC) detection channels have to be set manually by the user according to the signals of the controls.

1. In the Data Analysis window ensure to select the **Well Group** of the AltoStar® HAV RT-PCR Kit 1.5. Therefore, click on the **Well Group** drop-down menu right next to the **Well Group** button (see figure 4) of the toolbar.

2. At the left side of the Data Analysis window tick only the **VIC** checkbox for the detection channel of the IC (see figure 9).

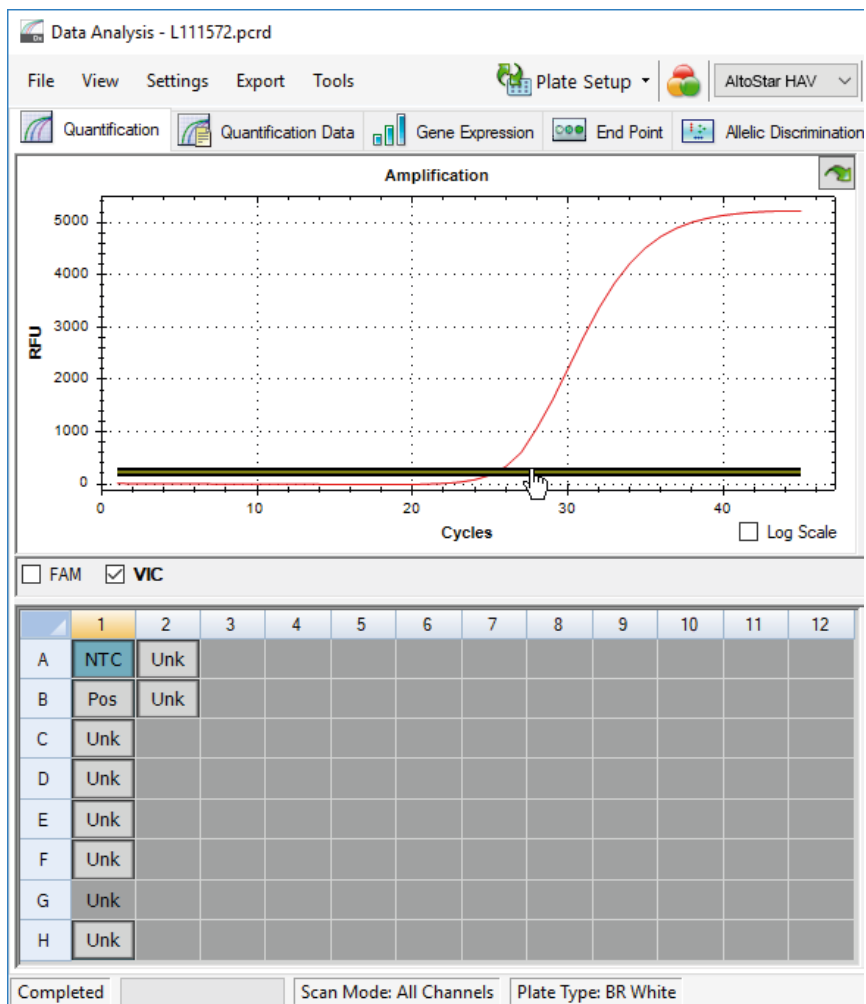


Figure 9: Data Analysis window: setting the VIC™ threshold

3. Select only the NTC well in the plate view of the Data Analysis window (see figure 9).
4. Drag the threshold into the exponential area of the NTC signal (see figure 9).

NOTE



The NTC contains the IC template, which leads to an IC signal in a valid NTC well.

5. At the left side of the Data Analysis window untick the **VIC** checkbox and tick the **FAM** checkbox for the detection channel of the HAV target (see figure 10).

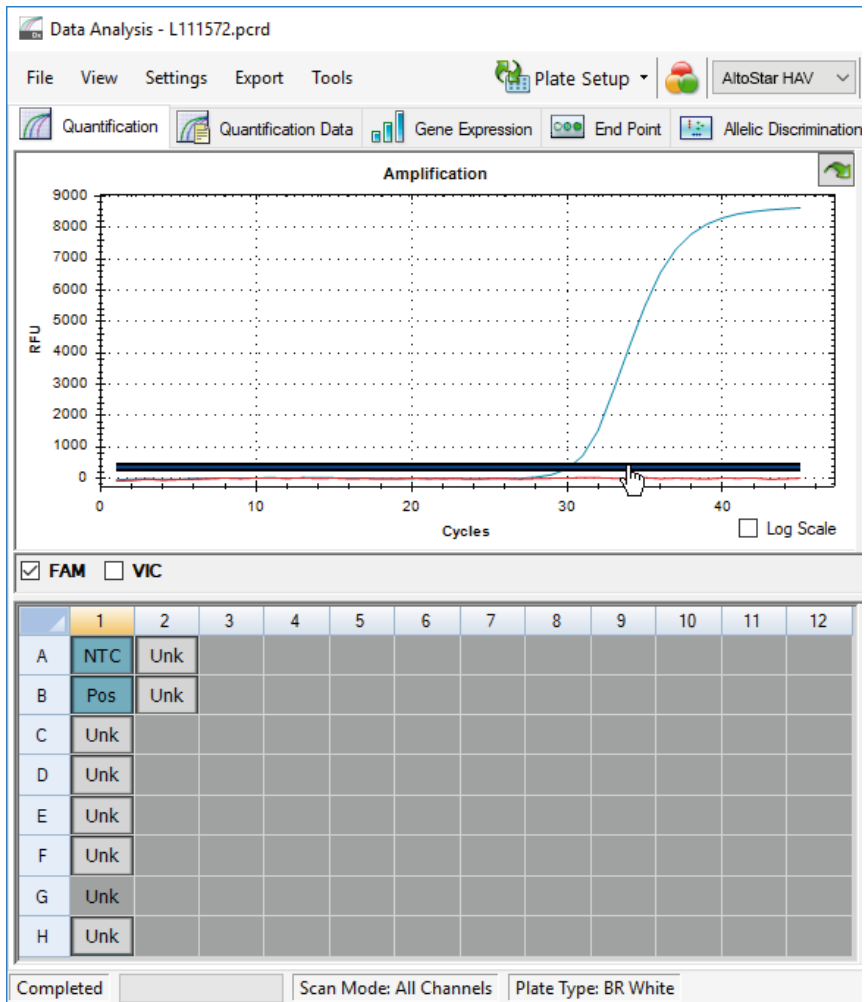


Figure 10: Data Analysis window: setting the FAM™ threshold

6. Select only the wells containing the NTC and the PC in the plate view of the Data Analysis window (see figure 10).
7. Drag the threshold well above the signal of the NTC into the exponential area of the PC signal (see figure 10).

7.6.8 Validity of PCR results

7.6.8.1 Exclusion of wells containing invalid data

Wells that do not contain valid data have to be excluded from result generation by the user.

1. In the Data Analysis window ensure to select the **Well Group** of the AltoStar® HAV RT-PCR Kit 1.5. Therefore, click on the **Well Group** drop-down menu right next to the **Well Group** button (see figure 4) of the toolbar.
2. Identify all wells containing invalid data. A well is invalid if any of the following conditions apply:
 - a) The complete run is invalid (see chapter 7.6.8.2 Validity of a diagnostic PCR run).
 - b) The well data does not meet the control conditions for a valid result (see chapter 7.6.8.3 Validity of results for a sample).

- Click each well containing invalid data according to chapters 7.6.8.2 Validity of a diagnostic PCR run to 7.6.8.3 Validity of results for a sample with the right mouse button and select **Well...** → **Exclude from Analysis** (see figures 11 and 12).

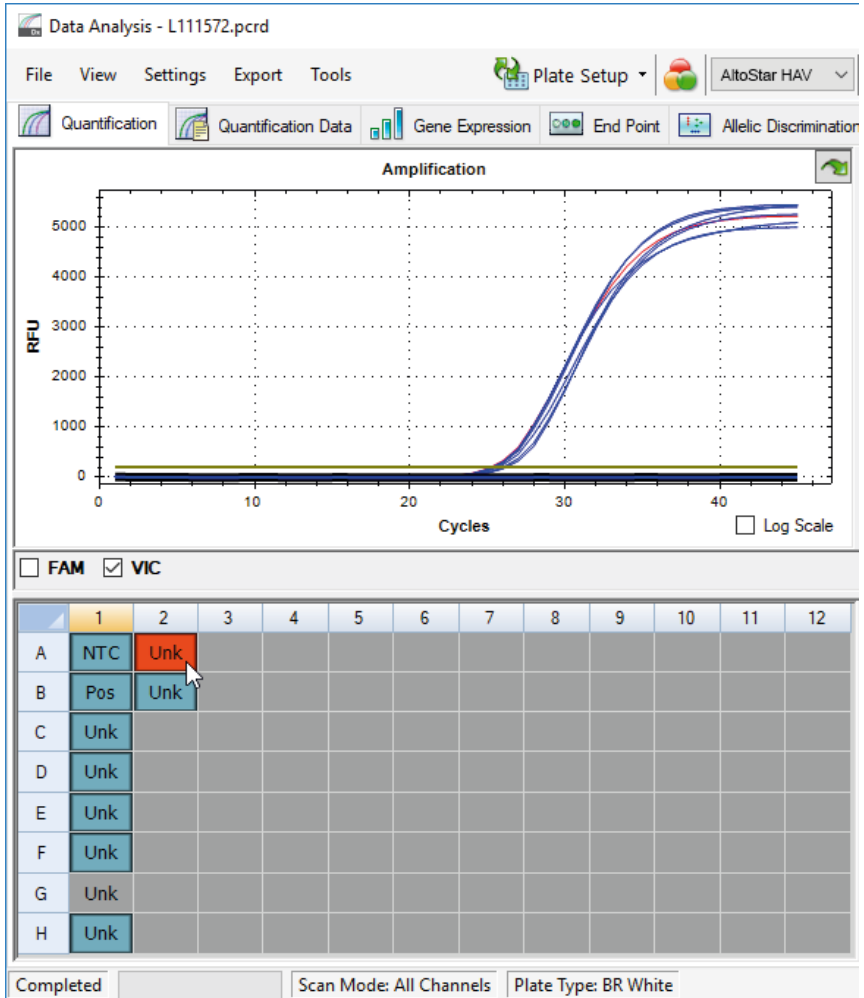


Figure 11: Data Analysis window: invalid well

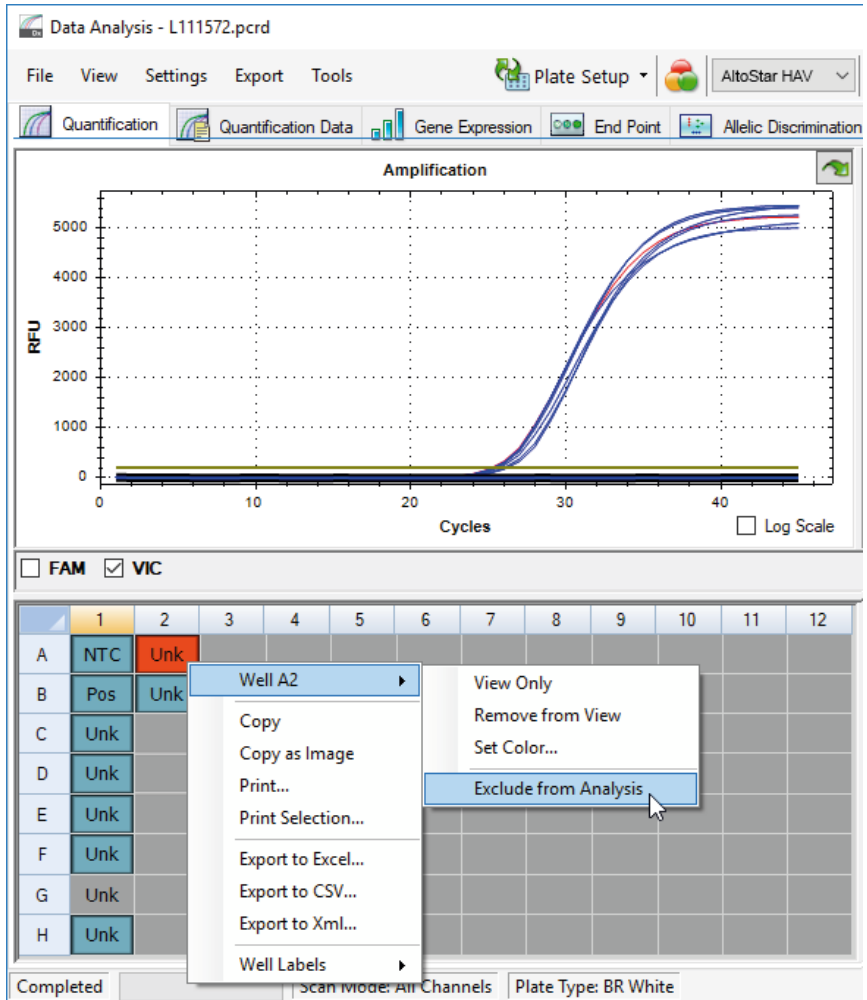


Figure 12: Data Analysis window: exclude invalid well from analysis

The selected well is excluded from the analysis. No results will be generated for this well.

7.6.8.2 Validity of a diagnostic PCR run

A diagnostic PCR run is **valid**, if the following control conditions are met:

Table 7: Control conditions for a valid PCR run

Control	Detection channel	
	FAM™ (HAV target)	VIC™ (IC)
PC	+	Not applicable
NTC	-	+

A diagnostic PCR run is **invalid**, if:

- The run has not been completed.
- Any of the control conditions for a valid diagnostic PCR run are not met.

In case of an invalid diagnostic PCR run, exclude all wells from the analysis and repeat the AltoStar® run starting from the original samples.

7.6.8.3 Validity of results for a sample

The result for an individual sample is **invalid**, if the signals in both the VIC™ (IC) detection channel and the FAM™ (HAV target) detection channel are negative (see table 8). In case of an invalid result for a sample, exclude the well from the analysis and repeat testing from the original sample or collect and test a new sample.

Table 8: Result validity

Detection channel		Result validity
FAM™ (HAV target)	VIC™ (IC)	
+	+	Valid result
+	-	Valid result*
-	+	Valid result
-	-	Invalid result

* Detection of the IC is not required when the HAV target is detected. A high HAV RNA load in the sample can lead to a reduced or absent IC signal.

7.6.9 Export of PCR results for result interpretation performed by LIMS

To make the PCR run results available to a connected LIMS for result interpretation performed by LIMS they need to be exported in the form of a LIMS result file (.CSV).

1. In the Data Analysis window ensure to select the **Well Group** of the AltoStar® HAV RT-PCR Kit 1.5. Therefore, click on the **Well Group** drop-down menu right next to the **Well Group** button (see figure 4) of the toolbar.
2. Ensure that all steps of the analysis process (see chapters 7.6.7.2 Baseline correction to 7.6.8.1 Exclusion of wells containing invalid data) have been completed for the well group of the AltoStar® HAV RT-PCR Kit 1.5.
3. In the menu bar of the Data Analysis window click **Export** → **Export All Data Sheets** to open the Browse For Folder dialog.
4. In the Browse For Folder dialog specify the location for the LIMS result files to be generated and click **OK**.

NOTE



The LIMS integration has to be implemented according to the specifications of altona Diagnostics. For information regarding LIMS integration, see chapter 17. Assay protocol for the AltoStar® Connect software and information for LIMS integration and/or contact altona Diagnostics technical support (see chapter 12. Technical support).

NOTE

Saving results of more than one assay (well group) from a PCR run in the same folder leads to replacement of the LIMS result files of the first assay (well group) with the LIMS result files of the second assay (well group). In this case the LIMS result files of the first assay (well group) can be exported again.

7.6.10 Export of PCR results for manual result interpretation

If the results are not passed to a LIMS for result interpretation performed by LIMS, the result interpretation has to be performed manually by the user. To this end the analysis results of each assay (well group) need to be exported in the form of a report.

1. In the Data Analysis window ensure to select the **Well Group** of the AltoStar® HAV RT-PCR Kit 1.5. Therefore, click on the **Well Group** drop-down menu right next to the **Well Group** button (see figure 4) of the toolbar.
2. At the left side of the Data Analysis window tick the **VIC** checkbox as well as the **FAM** checkbox.
3. Ensure that all steps of the analysis process (see chapters 7.6.7.2 Baseline correction to 7.6.8.1 Exclusion of wells containing invalid data) have been completed for the well group of the AltoStar® HAV RT-PCR Kit 1.5.
4. In the menu bar of the Data Analysis window click **Tools** → **Reports...** to open the Report dialog.

5. Ensure that at least the following content is selected for report generation in the upper left part of the Report dialog (see figure 13):

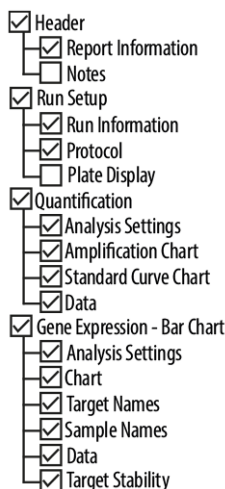


Figure 13: Report dialog

6. Select or deselect additional content of the report by ticking the respective check boxes as required.
7. In the menu bar of the Report dialog click **File** → **Save As...** to open the Save Report dialog.
8. In the Save Report dialog specify the name and location for the report file to be generated and click **Save**.

7.6.10.1 Manual interpretation of results

1. Open the report file generated for the well group of the AltoStar® HAV RT-PCR Kit 1.5 (see chapter 7.6.10 Export of PCR results for manual result interpretation).
2. Refer to the Quantification Data table in the report (see figure 14). The table comprises 2 rows for each **Sample** – one for the **Target HAV** and one for the **Target Internal Control**.

Quantification Data

Well	Fluor	Target	Content	Sample	Biological Set Name	Cq	Cq Mean	Cq Std. Dev	Well Note
A01	FAM	HAV	NTC	NTC 117201022202	AltoStar HAV RT-PCR Kit 1.5	N/A	0.00	0.000	
A02	FAM	HAV	Unkn	Sample 7 00000007	AltoStar HAV RT-PCR Kit 1.5	35.44	35.44	0.000	qualitative
B01	FAM	HAV	Pos Ctrl	HAV PC 117111022206	AltoStar HAV RT-PCR Kit 1.5	31.37	31.37	0.000	
B02	FAM	HAV	Unkn	Sample 8 00000008	AltoStar HAV RT-PCR Kit 1.5	35.08	35.08	0.000	qualitative
C01	FAM	HAV	Unkn	Sample 1 00000001	AltoStar HAV RT-PCR Kit 1.5	36.33	36.33	0.000	qualitative
D01	FAM	HAV	Unkn	Sample 2 00000002	AltoStar HAV RT-PCR Kit 1.5	36.57	36.57	0.000	qualitative
E01	FAM	HAV	Unkn	Sample 3 00000003	AltoStar HAV RT-PCR Kit 1.5	34.26	34.26	0.000	qualitative
F01	FAM	HAV	Unkn	Sample 4 00000004	AltoStar HAV RT-PCR Kit 1.5	35.39	35.39	0.000	qualitative
G01	FAM	HAV	Unkn	Sample 5 00000005	AltoStar HAV RT-PCR Kit 1.5	35.96	35.96	0.000	qualitative
H01	FAM	HAV	Unkn	Sample 6 00000006	AltoStar HAV RT-PCR Kit 1.5	34.93	34.93	0.000	qualitative
A01	VIC	Internal Control	NTC	NTC 117201022202	AltoStar HAV RT-PCR Kit 1.5	28.77	28.77	0.000	
A02	VIC	Internal Control	Unkn	Sample 7 00000007	AltoStar HAV RT-PCR Kit 1.5	29.43	29.43	0.000	qualitative
B02	VIC	Internal Control	Unkn	Sample 8 00000008	AltoStar HAV RT-PCR Kit 1.5	29.54	29.54	0.000	qualitative
C01	VIC	Internal Control	Unkn	Sample 1 00000001	AltoStar HAV RT-PCR Kit 1.5	29.22	29.22	0.000	qualitative
D01	VIC	Internal Control	Unkn	Sample 2 00000002	AltoStar HAV RT-PCR Kit 1.5	29.49	29.49	0.000	qualitative
E01	VIC	Internal Control	Unkn	Sample 3 00000003	AltoStar HAV RT-PCR Kit 1.5	29.50	29.50	0.000	qualitative
F01	VIC	Internal Control	Unkn	Sample 4 00000004	AltoStar HAV RT-PCR Kit 1.5	29.49	29.49	0.000	qualitative
G01	VIC	Internal Control	Unkn	Sample 5 00000005	AltoStar HAV RT-PCR Kit 1.5	29.27	29.27	0.000	qualitative
H01	VIC	Internal Control	Unkn	Sample 6 00000006	AltoStar HAV RT-PCR Kit 1.5	29.67	29.67	0.000	qualitative

Figure 14: Report: Quantification Data

Qualitative results would be marked by the term *qualitative* in the **Well Note** column of the Quantification Data table.

- Identify each row with the **Target** *HAV* and the term *qualitative* in the **Well Note** column.
- In these rows, refer to the **C_q** column for the result of the respective **Sample**.
- Refer to table 9 for interpretation of results.

Table 9: Result interpretation

Threshold cycle (C _q) of the HAV target	Result interpretation
1–45	HAV specific RNA detected.
N/A	No HAV specific RNA detected. Sample does not contain detectable amounts of HAV specific RNA.

7.6.11 PCR data analysis and automated result interpretation using FastFinder

Optionally, PCR data analysis and automated result interpretation can be done by using FastFinder Standalone and the AltoStar® HAV RT-PCR Kit 1.5 assay plugin. The valid assay plugin version is listed in chapter 3.2 AltoStar® HAV RT-PCR Kit 1.5 assay plugin.

For more information on FastFinder Standalone, refer to the respective instructions for use.

NOTE



The AltoStar® HAV RT-PCR Kit 1.5 assay plugin is intended to be used as support for the interpretation of the results generated with the AltoStar® HAV RT-PCR Kit 1.5 on the CFX96™ DW Dx. All results generated with the assay plugin have to be checked for validity and correctness by a human operator.

1. Start the FastFinder Standalone application on the AltoStar® Automated Analysis PC.
2. Enter your user data and password to log in. The start screen is displayed. All open and authorized analyses can be accessed.
3. Click on **Create new analysis**.
4. Browse for the data file to be analyzed and select it by clicking on **Open**. Only .PCRD files are supported. The file is uploaded and analyzed. After the analysis is finished **Ready for review** is displayed.

NOTE



It is highly recommended to analyze only one PCR data file (.PCRD) at once instead of analyzing multiple PCR runs in batches.

5. Click on the PCR run to start the review process.
6. Check controls and resolve any error messages based on the warning for individual samples.

NOTE

Make sure to confirm unresolvable samples as inconclusive and to re-extract and re-test the samples.

7. Authorize or restart the analysis.

After clicking on **Authorize** the analysis is automatically downloaded as .CSV file. PDFs can be downloaded from authorized analyses. Click the **Actions** button in the upper right corner and choose **Generate PDF Export**. Go to **Downloads** via the same menu to open the file.

8. Using the AltoStar® HAV RT-PCR Kit 1.5 with other real-time PCR instruments than the CFX96™ Deep Well Dx System

In addition to the CFX96™ DW Dx the AltoStar® HAV RT-PCR Kit 1.5 has been validated with other real-time PCR instruments (see chapter 5.3.2 Other workflows). The following chapters 8.1 Material and devices required but not provided and 8.2 Procedure describe how to use the AltoStar® HAV RT-PCR Kit 1.5 with these instruments.

8.1 Material and devices required but not provided

The following instruments, software and materials are required:

- General materials and devices (see chapter 7.5 General material and devices)
- CFX96™ Deep Well Dx System and the CFX Manager™ Dx software version 3.1 (Bio-Rad)
 - 96-well PCR plates and sealing foil (details see table 2)
- CFX96™ Dx System and the CFX Manager™ Dx software version 3.1 (Bio-Rad)
 - 96-well PCR plates and sealing foil (details see table 2)

- Rotor-Gene® Q5/6 plex Platform and the Rotor-Gene® Q Software version 2.3.1 (QIAGEN)
 - 0.1 ml strip tubes and lids [STRIP Tubes 0.1 ml for Rotor-Gene® cyclers (LTF Labortechnik) or equivalent]
- QuantStudio™ 5 Real-Time PCR System and the QuantStudio™ 5 Dx Software v1.0.2 and ABI Prism® 7500 SDS and the 7500 Software v2.3 (Applied Biosystems)
 - 96-well PCR plates and sealing foil [MicroAmp™ Optical 96-Well Reaction Plate and MicroAmp™ Optical Adhesive Film (Applied Biosystems) or equivalent]
- LightCycler® 480 Instrument II and the LightCycler® 480 Software version 1.5.1 (Roche)
 - 96-well PCR plates and sealing foil [LightCycler® 480 Multiwell Plate 96, white and LightCycler® 480 Sealing Foil (Roche) or equivalent]

NOTE



It is not recommended to use other materials or devices than specified in these instructions for use.

8.2 Procedure

8.2.1 Sample preparation

The AltoStar® HAV RT-PCR Kit 1.5 was validated using the AltoStar® AM16 in combination with the AltoStar® Purification Kit 1.5.

The AltoStar® HAV RT-PCR Kit 1.5 has to be used with a heterologous IC (AltoStar® Internal Control 1.5), which allows to control the sample preparation procedure (nucleic acid extraction) and the following RT-PCR.

The volume of the IC which is added to the sample/lysis buffer mixture depends on the elution volume. It represents 50 % of the elution volume.

CAUTION

Storage of eluates under wrong conditions may lead to degradation of the HAV target sequence and could compromise product performance.

8.2.2 Master mix setup

All components of the AltoStar® HAV RT-PCR Kit 1.5 and samples should be thawed completely, mixed (by pipetting or gentle vortexing) and centrifuged briefly before use. Set up the master mix according to the following pipetting scheme:

Table 10: Pipetting scheme (master mix setup)

Number of reactions (rxns)	1	12
Master A	5 µl	60 µl
Master B	15 µl	180 µl
Volume master mix	20 µl	240 µl

CAUTION

Do not use different volumes of Master A and Master B for master mix setup than specified in these instructions for use, as this could compromise product performance.

CAUTION

A lack of centrifugation of the product components after thawing may cause contamination with reagent residues in the lids and could compromise product performance.

8.2.3 Reaction setup

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

Table 11: Pipetting scheme (reaction setup)

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

3. Make sure that at least 1 PC and 1 NTC is used per run.
4. Thoroughly mix the samples and controls with the master mix by pipetting up and down.
5. Close the 96-well PCR plate with a PCR plate sealing foil and the reaction tubes with appropriate lids (see chapter 8.1 Material and devices required but not provided).
6. Centrifuge the 96-well PCR plate in a centrifuge with a microtiter plate rotor for 30 seconds at approximately 1,000 x g (~ 3,000 rpm).

The NTC does always contain the IC template in the correct concentration.

After completion of the PCR setup, the PCR mix is stable at room temperature (max. +30 °C) for 30 minutes.

CAUTION



Do not exceed the PCR mix storage time, as this could compromise product performance.

CAUTION

Do not mix up samples or sample IDs during PCR setup or transfer to the PCR instrument. This could lead to false positive or false negative results due to incorrect assignment of the samples.

8.2.4 PCR run

8.2.4.1 Programming the real-time PCR instrument

For basic information regarding the setup and programming of the different real-time PCR instruments, refer to the instructions for use of the respective instrument.

For detailed programming instructions regarding the use of the AltoStar® HAV RT-PCR Kit 1.5 on specific real-time PCR instruments, contact Altona Diagnostics technical support (see chapter 12. Technical support).

8.2.4.2 Run settings

Define the following basic settings:

Table 12: Run settings

Settings	
Reaction volume	30 µl
Ramp rate	Default
Passive reference*	ROX™

* If applicable

Define the following fluorescence detectors (dyes):

Table 13: Fluorescence detectors

Target	Detector name	Reporter	Quencher
HAV	HAV	FAM™	(None)
IC	Internal Control	JOE™	(None)

Define the following temperature profile and dye acquisition:

Table 14: Temperature profile and dye acquisition

	Stage	Cycle repeats	Acquisition	Temperature [°C]	Time [min:s]
Reverse transcription	Hold	1	-	55	20:00
Denaturation	Hold	1	-	95	02:00
Amplification	Cycling	45	-	95	00:15
			Yes	55	00:45
			-	72	00:15

CAUTION



Do not use other cycling conditions than specified in these instructions for use, as this could compromise product performance.

8.2.5 Data analysis

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

For detailed instructions regarding the analysis of the data generated with the AltoStar® HAV RT-PCR Kit 1.5 on different real-time PCR instruments, contact altona Diagnostics technical support (see chapter 12. Technical support).

Criteria for validity of diagnostic PCR runs and result interpretation regardless of the real-time PCR instrument are described in chapters 7.6.8.2 Validity of a diagnostic PCR run and 7.6.8.3 Validity of results for a sample, chapter 7.6.10.1 Manual interpretation of results and table 9.

CAUTION



Do not use other control settings for data analysis than specified in these instructions for use, as this could cause incorrect IVD examination results.

CAUTION



As with any diagnostic test, results shall be interpreted in consideration of all clinical and laboratory findings.

9. Performance data

The performance of the AltoStar® HAV RT-PCR Kit 1.5 was evaluated using the 2nd and 3rd WHO International Standard for hepatitis A virus for Nucleic Acid Amplification Techniques (NIBSC code: 00/562 and 15/276, respectively) and HAV standard material calibrated against the WHO International Standard.

The performance data were obtained with the AltoStar® HAV RT-PCR Kit 1.5 using the AltoStar® Workflow. Equivalent performance in terms of specificity, sensitivity and precision was demonstrated for the other real-time PCR instruments listed in chapter 5.3.2 Other workflows.

9.1 Plasma

9.1.1 Analytical sensitivity

For the determination of the limit of detection (LoD) a dilution series of the 3rd WHO International Standard for hepatitis A virus for Nucleic Acid Amplification Techniques (NIBSC code: 15/276) in plasma ranging from 1.00E+02 to 2.50E-01 IU/ml was generated.

Each dilution was tested in 8 replicates in 3 different runs (total n = 24 per dilution) using combinations of:

- 3 AltoStar® HAV RT-PCR Kit 1.5 lots
- 3 AltoStar® Purification Kit 1.5 lots
- 3 AltoStar® Internal Control 1.5 lots
- 3 AltoStar® AM16 instruments
- 3 CFX96™ DW Dx instruments

Data from all runs were combined and a probit analysis was performed to determine the 95 % LoD value.

Table 15: PCR results used for the calculation of the analytical sensitivity of the AltoStar® HAV RT-PCR Kit 1.5

Concentration [IU/ml]	N [total]	N [positive]	Hit rate [%]
1.00E+02	24	24	100
5.00E+01	24	24	100
2.50E+01	24	24	100
1.00E+01	24	24	100
5.00E+00	24	23	96
2.50E+00	24	16	67
1.00E+00	24	12	50
5.00E-01	24	4	17
2.50E-01	24	3	13

The LoD of the AltoStar® HAV RT-PCR Kit 1.5 for the detection of HAV in plasma is 6.31 IU/ml (95 % confidence interval: 4.17–11.99 IU/ml).

9.1.2 Analytical specificity

The analytical specificity of the AltoStar® HAV RT-PCR Kit 1.5 is ensured by the thorough selection of the oligonucleotides (primers and probes). The oligonucleotides were checked by sequence comparison analysis against publicly available sequences to ensure that all relevant HAV genotypes will be detected.

For the verification of the analytical specificity of the AltoStar® HAV RT-PCR Kit 1.5 the following experiments were performed (see chapters 9.1.2.1 Negative samples to 9.1.2.3 Cross-reactivity).

9.1.2.1 Negative samples

35 HAV negative plasma samples from individual donors were tested with the AltoStar® HAV RT-PCR Kit 1.5. All (35 out of 35) samples were tested negative for HAV specific RNA and positive for the IC. The analytical specificity of the AltoStar® HAV RT-PCR Kit 1.5 for plasma samples is $\geq 95\%$.

9.1.2.2 Interfering substances

To evaluate the influence of potentially interfering endogenous substances on the performance of the AltoStar® HAV RT-PCR Kit 1.5 selected substances were spiked in plasma samples. These plasma samples contained HAV in a concentration of the 3 x LoD ($1.90E+01$ IU/ml) and no HAV, respectively.

Results obtained for samples containing potentially interfering substances were compared to results generated for plasma samples containing no spiked interferent. Each sample was processed in 3 replicates.

No interference was observed for samples containing elevated levels of:

- Endogenous substances
 - Bilirubin
 - Hemoglobin
 - Human genomic DNA
 - Human serum albumin
 - Triglycerides

CAUTION



The presence of PCR inhibitors (e.g., heparin) could cause false negative or invalid results.

9.1.2.3 Cross-reactivity

The analytical specificity of the AltoStar® HAV RT-PCR Kit 1.5 with respect to cross-reactivity with other pathogens than HAV was evaluated by testing:

- Viruses related to HAV
- Viruses causing similar symptoms as an infection with HAV
- Viruses likely to be present in patients suffering from a HAV infection

The AltoStar® HAV RT-PCR Kit 1.5 did not cross-react with any of the following pathogens:

- Coxsackievirus A3
- Cytomegalovirus (CMV)
- Enterovirus 71
- Epstein-Barr virus (EBV)
- Hepatitis B virus (HBV)
- Hepatitis C virus (HCV)
- Hepatitis E virus (HEV)
- Herpes simplex virus 1 (HSV-1)
- Herpes simplex virus 2 (HSV-2)
- Human immunodeficiency virus 1 (HIV-1)
- Parvovirus B19

- Rubella virus
- Rubulavirus (RuV)
- Varicella-zoster virus (VZV)
- Yellow fever virus

CAUTION



In case the sample contains other pathogens than HAV, competition with the target amplification or cross-reactivities may occur, causing incorrect IVD examination results.

9.1.3 Reactivity

Specificity of the AltoStar® HAV RT-PCR Kit 1.5 with respect to the detection of HAV genotypes is foremost ensured by the selection of the primers and probes. To verify that the AltoStar® HAV RT-PCR Kit 1.5 allows the detection of other HAV genotypes than HAV 1a [2nd WHO International Standard for hepatitis A virus (NIBSC code: 00/562)] and 1b [3rd WHO International Standard for hepatitis A virus (NIBSC code: 15/276)], HAV 2a und 3a were tested with the AltoStar® HAV RT-PCR Kit 1.5. HAV 2a and 3a were detected positive with the AltoStar® HAV RT-PCR Kit 1.5.

CAUTION



Potential mutations within the target regions of the HAV genome covered by primers and/or probes used in the kit may result in failure to detect the presence of the pathogen.

9.1.4 Precision

Precision of the AltoStar® HAV RT-PCR Kit 1.5 was evaluated using a panel consisting of:

- 1 HAV high positive plasma sample (3.15E+02 IU/ml)
- 1 HAV low positive plasma sample [1.90E+01 IU/ml (3 x LoD)]
- 1 HAV negative plasma sample

Data were determined based on:

- Threshold cycle (C_q^*) values for the HAV high positive sample
- Threshold cycle (C_q^*) values for the IC in the HAV negative sample

* Please note that the chosen term C_q is equivalent to the designation of C_t , which might be used by other cyclers than the CFX96™ Deep Well Dx System (Bio-Rad).

Repeatability data were generated following recommendations of the CLSI guideline EP05-A3 (“Evaluation of Precision of Quantitative Measurement Procedures; Approved Guideline—Third Edition”) [3]. Each panel member was tested in duplicates in 2 runs per day over 20 days by one operator using:

- 1 AltoStar® HAV RT-PCR Kit 1.5 lot
- 1 AltoStar® Purification Kit 1.5 lot
- 1 AltoStar® Internal Control 1.5 lot
- 1 AltoStar® AM16 instrument
- 1 CFX96™ DW Dx instrument

In table 16 the estimates of repeatability and within-laboratory precision are provided both as standard deviation (SD) and as coefficient of variation (CV %) based on C_q values obtained for the HAV high positive sample and for the IC in the HAV negative sample. The precision data shown were obtained by analyzing the data manually with the CFX Manager™ Dx software and using the automated result interpretation of FastFinder Standalone and the AltoStar® HAV RT-PCR Kit 1.5 assay plugin.

Table 16: Repeatability and within-laboratory precision for the AltoStar® HAV RT-PCR Kit 1.5

		Repeatability			Within-laboratory precision	
CFX Manager™ Dx software						
Sample	Mean (C _q values)	SD (C _q values)	CV %	SD (C _q values)	CV %	
High positive	30.27	0.11	0.36	0.13	0.42	
Negative	29.37	0.10	0.35	0.14	0.47	
AltoStar® HAV RT-PCR Kit 1.5 assay plugin						
Sample	Mean (C _q values)	SD (C _q values)	CV %	SD (C _q values)	CV %	
High positive	28.29	0.12	0.44	0.32	1.12	
Negative	27.84	0.13	0.46	0.36	1.30	

All samples tested at 3 x LoD (low positive samples) were detected positive and all negative samples resulted negative with the CFX Manager™ Dx software and the AltoStar® HAV RT-PCR Kit 1.5 assay plugin.

For the reproducibility, each panel member was tested in 6 replicates in 3 runs by 3 operators using:

- 3 AltoStar® HAV RT-PCR Kit 1.5 lots
- 3 AltoStar® Purification Kit 1.5 lots
- 3 AltoStar® Internal Control 1.5 lots
- 3 AltoStar® AM16 instruments
- 3 CFX96™ DW Dx instruments

In table 17 the estimates of reproducibility are provided both as standard deviation (SD) and as coefficient of variation (CV %) based on C_q obtained for the HAV high positive sample and the IC in the HAV negative sample. The precision data shown were obtained by analyzing the data manually with the CFX Manager™ Dx software and using the automated result interpretation of FastFinder Standalone and the AltoStar® HAV RT-PCR Kit 1.5 assay plugin.

Table 17: Reproducibility for the AltoStar® HAV RT-PCR Kit 1.5

		Reproducibility	
CFX Manager™ Dx software			
Sample	Mean (C_q values)	SD (C_q values)	CV %
High positive	30.53	0.26	0.85
Negative	29.57	0.19	0.66
AltoStar® HAV RT-PCR Kit 1.5 assay plugin			
Sample	Mean (C_q values)	SD (C_q values)	CV %
High positive	28.37	0.17	0.61
Negative	27.30	0.17	0.63

All samples tested at 3 x LoD (low positive samples) were detected positive and all negative samples resulted negative with the CFX Manager™ Dx software and the AltoStar® HAV RT-PCR Kit 1.5 assay plugin.

9.1.5 Total failure rate

The robustness of the AltoStar® HAV RT-PCR Kit 1.5 was assessed by testing 33 HAV negative plasma samples from individual donors spiked with HAV to a final concentration of the 3 x LoD (1.90E+01 IU/ml). All (33 out of 33) samples were tested positive in the HAV specific fluorescence detection channel (FAM™).

9.1.6 Carry over

Carry over is mostly a workflow dependent risk and independent of the PCR assay used. For the AltoStar® Workflow the AltoStar® Parvovirus B19 PCR Kit 1.5 was used as exemplary model. Potential cross-contamination through carry over from high positive samples was evaluated by testing alternating parvovirus B19 high positive ($1.00E+07$ IU/ml) and negative samples ($n = 44$ each per run; 5 runs) with the AltoStar® Parvovirus B19 PCR Kit 1.5. No carry over was observed, i.e. all parvovirus B19 negative samples were tested negative.

9.1.7 Clinical performance

The AltoStar® HAV RT-PCR Kit 1.5 was evaluated in a comparative study with the CE-marked AmpliSens® HAV-FRT PCR Kit (AmpliSens Ecoli Dx, s.r.o.). Retrospectively, 145 human plasma samples were tested in parallel:

The AmpliSens® HAV-FRT PCR Kit (AmpliSens Ecoli Dx, s.r.o.) was used in combination with the NucliSENS® easyMAG instrument (bioMérieux) and the RotorGene® 6000 instrument (QIAGEN), respectively the Mx3000P instrument (Agilent).

The AltoStar® HAV RT-PCR Kit 1.5 was used in combination with the AltoStar® Purification Kit 1.5 and the AltoStar® Internal Control 1.5 on the AltoStar® AM16 and the CFX96™ DW Dx.

For the qualitative analysis all samples with an invalid result for one or both assays were excluded.

Results for the remaining 139 samples are shown in table 18.

Table 18: Results of the evaluation of the diagnostic sensitivity and specificity for HAV in plasma samples

		AmpliSens® HAV-FRT PCR Kit (AmpliSens Ecoli Dx, s.r.o.)	
		POSITIVE	NEGATIVE
AltoStar® HAV RT-PCR Kit 1.5	POSITIVE	68	2
	NEGATIVE	0	69

The diagnostic sensitivity and specificity of the AltoStar® HAV RT-PCR Kit 1.5 compared to the AmpliSens® HAV-FRT PCR Kit (AmpliSens Ecoli Dx, s.r.o.) were 100 % and 97 %, respectively.

10. Disposal

Dispose of hazardous and biological waste in compliance with local and national regulations. Leftover product components and waste should not be allowed to enter sewage, water courses or the soil.

CAUTION



Always treat samples as infectious and (bio-)hazardous material in accordance with safety and laboratory procedures. For sample material spills promptly use an appropriate disinfectant. Handle contaminated materials as biohazardous.

CAUTION



Disposal of hazardous and biological waste shall comply with local and national regulations to avoid environmental contamination.

NOTE



The PCR plate must be disposed of in a sealed state since the PCR plate sealing foil cannot be removed.

11. Quality control

In accordance with the Altona Diagnostics GmbH EN ISO 13485-certified Quality Management System, each lot of AltoStar® HAV RT-PCR Kit 1.5 is tested against predetermined specifications to ensure consistent product quality.

12. Technical support

For customer support, contact Altona Diagnostics technical support:

e-mail: support@altona-diagnostics.com

phone: +49-(0)40-5480676-0

NOTE



Any serious incident that has occurred in relation to this product shall be reported to Altona Diagnostics and the competent authority of your country.

13. Summary of safety and performance

The summary of safety and performance drawn up for the AltoStar® HAV RT-PCR Kit 1.5 can be found in the European Database on Medical Devices (EUDAMED): <https://ec.europa.eu/tools/eudamed>.

14. Literature

- [1] Versalovic, James, Carroll, Karen C., Funke, Guido, Jorgensen, James H., Landry, Marie Louise and David W. Warnock (ed). *Manual of Clinical Microbiology*. 10th Edition. ASM Press, 2011.
- [2] Cohen, Jonathan, Powderly, William G, and Steven M Opal. *Infectious Diseases*, Third Edition. Mosby, 2010.
- [3] CLSI. *Evaluation of Precision of Quantitative Measurement Procedures; Approved Guideline—Third Edition*. CLSI document EP05-A3. Wayne, PA: Clinical and Laboratory Standards Institute; 2014.

15. Trademarks and disclaimers

4s3™ (4titude); AltoStar® (altona Diagnostics); AmpliSens® (AmpliSens Ecoli Dx, s.r.o.); ABI Prism®, QuantStudio™ (Applied Biosystems); NucliSENS® (bioMérieux); CFX96™, CFX Manager™ (Bio-Rad); Rotor-Gene® (QIAGEN); LOINC® (Regenstrief Institute, Inc.); LightCycler® (Roche); FAM™, JOE™, MicroAmp™, VIC™ (Thermo Fisher Scientific).

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The AltoStar® HAV RT-PCR Kit 1.5 is a CE-marked product according to Regulation (EU) 2017/746 on *in vitro* diagnostic medical devices.









Product not licensed with Health Canada and not FDA cleared or approved.

Not available in all countries.

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16. Symbols

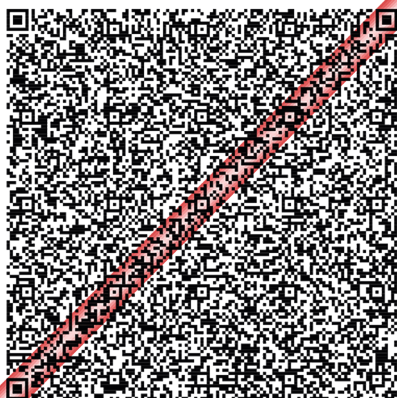
Symbol	Explanation
	<i>In vitro</i> diagnostic medical device
	Global Trade Item Number
	Batch code
	Content
	Cap color
	Catalogue number
	Number
	Component
	Material number
	Version number
	Version
	Consult instructions for use

Symbol	Explanation
	Contains sufficient for "n" tests/reactions (rxns)
	Temperature limit
	Use-by date
	Manufacturer
	Date of manufacture
	Caution
	Note
	Contains biological material of animal origin

17. Assay protocol for the AltoStar® Connect software and information for LIMS integration

The 2D barcode in figure 15 is to be used for installing the latest assay protocol for use of the AltoStar® HAV RT-PCR Kit 1.5 on the AltoStar® AM16. The barcode can only be scanned in printed form. You can scan the barcode directly from the manual or print it on a separate sheet. Please note that the size of the print affects the scannability of the barcode. Make sure to scale the size to 100 %. For scanning, aim the scanner to the red line on the barcode. For details on management of assay protocols, please refer to the respective chapter in the instructions for use of the AltoStar® Connect software. For information for LIMS integration, see table 20.

AltoStar® HAV RT-PCR Kit 1.5



Protocol Version:

1

Checksum: C326507A464335D02618D2794862824FFF75E7DB

Figure 15: Assay protocol barcode for the AltoStar® HAV RT-PCR Kit 1.5

Table 19: Changelog for the assay protocol

Protocol version	Release updates
1	Initial version

Table 20: Information for LIMS integration

Usage	Data
Test order (LIMS → AltoStar® AM16)	AltoStar HAV RT-PCR Kit 1.5
Test result (CFX96™ DW Dx → LIMS) unit	N/A
Test result (CFX96™ DW Dx → LIMS) channel 1	HAV
Test result (CFX96™ DW Dx → LIMS) channel 2	Internal Control

For LOINC® (Logical Observation Identifiers Names and Codes) please refer to the altona Diagnostics GmbH website (www.altona-diagnostics.com) or contact altona Diagnostics technical support (see chapter 12. Technical support).

18. Revision history

Table 21: Revision history

Identifier	Date of issue [month/year]	Modifications
MAN-AS0241540-EN-S05	12/2025	<ul style="list-style-type: none"> • Chapter 1: deletion of text section "nucleic acid extraction methods and" • Chapters 2, 5, 5.4.2: specification of sample types • Chapter 3: renaming chapter into "Kit components" • Chapter "3.1 Reagents" added containing the content from former chapter 3 • Chapter "3.2 AltoStar® HAV RT-PCR Kit 1.5 assay plugin" added • Chapter 5: addition of target region • Chapter 5.2: renaming chapter into "Reagent description" • Chapters 5.3.1, 7, 7.4, 7.6.1, 7.6.11: addition of FastFinder for data analysis and automated result interpretation (optional) • Chapter 5.3.2: addition of content from former chapter 5.3.2.2 • Chapters 5.3.2, 8.1: addition of software versions • Deletion of chapters 5.3.2.1 and 5.3.2.2 • Chapters 5.4.2, 6: addition of cautions • Chapter 8.2.1: deletion of description regarding how to proceed for other nucleic acid extraction methods than the AltoStar® AM16 • Chapter 9: addition of information regarding performance data on other real-time PCR instruments • Chapter 9.1.4: exchange of precision data based on 5 runs with precision data generated in accordance with CLSI Guideline EP05-A3

Identifier	Date of issue [month/year]	Modifications
MAN-AS0241540- EN-S05	12/2025	<ul style="list-style-type: none">• Chapter "13. Summary of safety and performance" added• Chapter 14: addition of reference for CLSI Guideline EP05-A3• Chapter 15: exchange of "European <i>in vitro</i> diagnostic directive 98/79/EC" with "Regulation (EU) 2017/746 on <i>in vitro</i> diagnostic medical devices"• Editorial changes

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