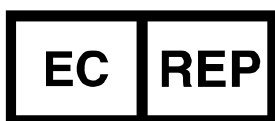




For Professional Use Only

**AmpliSens<sup>®</sup> *C.trachomatis* / *Ureaplasma* /  
*M.genitalium*-MULTIPRIME-FRT  
PCR kit  
Instruction Manual**

**AmpliSens<sup>®</sup>**



Ecoli s.r.o., Studenohorska 12  
841 03 Bratislava 47  
Slovak Republic  
Tel.: +421 2 6478 9336  
Fax: +421 2 6478 9040



Federal Budget Institute of  
Science "Central Research  
Institute for Epidemiology"  
3A Novogireevskaya Street  
Moscow 111123 Russia

## TABLE OF CONTENTS

1. INTENDED USE .....	3
2. PRINCIPLE OF PCR DETECTION .....	3
3. CONTENT .....	3
4. ADDITIONAL REQUIREMENTS .....	5
5. GENERAL PRECAUTIONS.....	5
6. SAMPLING AND HANDLING .....	5
7. WORKING CONDITIONS.....	6
8. PROTOCOL .....	6
9. DATA ANALYSIS .....	8
10. TROUBLESHOOTING.....	10
11. TRANSPORTATION.....	10
12. STABILITY AND STORAGE.....	10
13. SPECIFICATIONS.....	11
14. REFERENCES .....	11
15. QUALITY CONTROL.....	12
16. KEY TO SYMBOLS USED .....	13

## 1. INTENDED USE

**AmpliSens® C.trachomatis / Ureaplasma / M.genitalium-MULTIPRIME-FRT** PCR kit is an *in vitro* nucleic acid amplification test for simultaneous detection of DNA of *Chlamydia trachomatis*, *Ureaplasma* spp. (*U.parvum* and *U.urealyticum*), and *Mycoplasma genitalium* in the clinical material (urogenital, rectal and oropharyngeal swabs; conjunctival discharge; prostate gland secretion; and urine samples) by using real-time hybridization-fluorescence detection.



The results of PCR analysis are taken into account in complex diagnostics of disease.

## 2. PRINCIPLE OF PCR DETECTION

*Chlamydia trachomatis*, *Ureaplasma* spp. (*U.parvum* and *U.urealyticum*), *Mycoplasma genitalium* detection by the multiplex polymerase chain reaction (PCR) is based on the amplification of pathogen genome specific region using specific primers. In the real-time PCR, the amplified product is detected with the use of fluorescent dyes. These dyes are linked to oligonucleotide probes, which bind specifically to the amplified product during thermocycling. The real-time monitoring of fluorescence intensities during the real-time PCR allows the detection of accumulating product without re-opening the reaction tubes after the PCR run.

**AmpliSens® C.trachomatis / Ureaplasma / M.genitalium-MULTIPRIME-FRT** PCR kit is a qualitative test that contains the Internal Control (Internal Control-FL (IC)). It must be used in the extraction procedure in order to control the extraction process of each individual sample and to identify possible reaction inhibition.

**AmpliSens® C.trachomatis / Ureaplasma / M.genitalium-MULTIPRIME-FRT** PCR kit uses “hot-start”, which greatly reduces the frequency of nonspecifically primed reactions. In variant FRT, “hot-start” is guaranteed by the separation of nucleotides and Taq-polymerase by using a wax layer. Wax melts and reaction components mix only at 95 °C. In variant FRT-100 F, “hot-start” is guaranteed by the separation of nucleotides and Taq-polymerase by using chemically modified polymerase (TaqF). The chemically modified polymerase (TaqF) is activated by heating at 95 °C for 15 min.

## 3. CONTENT

**AmpliSens® C.trachomatis / Ureaplasma / M.genitalium-MULTIPRIME-FRT** PCR kit is produced in 2 forms:

**AmpliSens® C.trachomatis / Ureaplasma / M.genitalium-MULTIPRIME-FRT** PCR kit

variant FRT, **REF** R-B46-(RG)-CE.

**AmpliSens® C.trachomatis / Ureaplasma / M.genitalium-MULTIPRIME-FRT** PCR kit  
variant FRT-100 F, **REF** R-B46-F(RG,iQ)-CE.

**AmpliSens® C.trachomatis / Ureaplasma / M.genitalium-MULTIPRIME-FRT** PCR kit,  
variant FRT includes:

<b>Reagent</b>	<b>Description</b>	<b>Volume, ml</b>	<b>Quantity</b>
<b>PCR-mix-1-FL C.trachomatis / Ureaplasma / M.genitalium</b> ready-to-use single-dose test tubes ( <i>under wax</i> )	colorless clear liquid	0.01	110 tubes of 0.2 ml
<b>PCR-mix-2-FL-red</b>	red clear liquid	1.1	1 tube
<b>Positive Control complex (C+)</b>	colorless clear liquid	0.2	1 tube
<b>DNA-buffer</b>	colorless clear liquid	0.5	1 tube
<b>Negative Control (C-)*</b>	colorless clear liquid	1.2	1 tube
<b>Internal Control-FL (IC)**</b>	colorless clear liquid	1.0	1 tube

\* must be used in the extraction procedure as Negative Control of Extraction.

\*\* add 10 µl of Internal Control-FL (IC) during the DNA extraction procedure directly to the sample/lysis mixture (see DNA-sorb-AM **REF** K1-12-100-CE protocol).

**AmpliSens® C.trachomatis / Ureaplasma / M.genitalium-MULTIPRIME-FRT** PCR kit  
variant FRT is intended for 110 reactions, including controls.

**AmpliSens® C.trachomatis / Ureaplasma / M.genitalium-MULTIPRIME-FRT** PCR kit,  
variant FRT-100 F includes:

<b>Reagent</b>	<b>Description</b>	<b>Volume, ml</b>	<b>Quantity</b>
<b>PCR-mix-1-FL C.trachomatis / Ureaplasma / M.genitalium</b>	colorless clear liquid	1.2	1 tube
<b>PCR-mix-2-FRT</b>	colorless clear liquid	0.3	2 tubes
<b>Polymerase (TaqF)</b>	colorless clear liquid	0.03	2 tubes
<b>Positive Control complex (C+)</b>	colorless clear liquid	0.2	1 tube
<b>DNA-buffer</b>	colorless clear liquid	0.5	1 tube
<b>Negative Control (C-)*</b>	colorless clear liquid	1.2	1 tube
<b>Internal Control-FL (IC)**</b>	colorless clear liquid	1.0	1 tube

\* must be used in the extraction procedure as Negative Control of Extraction.

\*\* add 10 µl of Internal Control-FL (IC) during the DNA extraction procedure directly to the sample/lysis mixture (see “DNA-sorb-AM” **REF** K1-12-100-CE protocol).

**AmpliSens® C.trachomatis / Ureaplasma / M.genitalium-MULTIPRIME-FRT** PCR kit variant FRT-100 F is intended for 110 reactions, including controls.

#### 4. ADDITIONAL REQUIREMENTS

- Transport medium.
- DNA extraction kit.
- Disposable powder-free gloves and a laboratory coat.
- Pipettes (adjustable).
- Sterile pipette tips with filter (up to 200 µl).
- Tube racks.
- Vortex mixer.
- Desktop centrifuge with rotor for 2 ml reaction tubes.
- PCR box.
- Real-time instruments (for example, Rotor-Gene 3000/6000 (Corbett Research, Australia); Rotor-Gene Q (QIAGEN, Germany), iCycler iQ5 (Bio-Rad, USA); Mx3000P (Stratagene, USA)).
- Disposable polypropylene PCR tubes (0.1- or 0.2-ml) when working with PCR kit variant FRT-100 F:
  - a) 0.2-ml thin-walled PCR tubes with domed caps if a plate-type instrument is used;
  - b) 0.2-ml thin-walled PCR tubes with flat caps or strips of four 0.1-ml Rotor-Gene PCR tubes if a rotor-type instrument is used.
- Refrigerator for 2–8 °C.
- Deep-freezer for the temperature from minus 24 to minus 16 °C.
- Reservoir for used tips.

#### 5. GENERAL PRECAUTIONS

The user should always pay attention to the following:

- Use sterile pipette tips with aerosol barriers and use a new tip for every procedure.
- Store all extracted positive material (specimens, controls and amplicons) away from all other reagents and add it to the reaction mix in a distantly separated facility.
- Thaw all components thoroughly at room temperature before starting an assay.
- When thawed, mix the components and centrifuge briefly.
- Use disposable protective gloves and laboratory cloths, and protect eyes while samples

and reagents handling. Thoroughly wash hands afterwards.

- Do not eat, drink, smoke, apply cosmetics, or handle contact lenses in laboratory work areas.
- Do not use a kit after its expiration date.
- Dispose of all specimens and unused reagents in accordance with local regulations.
- Samples should be considered potentially infectious and handled in biological cabinet in compliance with appropriate biosafety practices.
- Clean and disinfect all samples or reagents spills using a disinfectant, such as 0.5 % sodium hypochlorite or another suitable disinfectant.
- Avoid samples and reagents contact with the skin, eyes, and mucous membranes. If these solutions come into contact, rinse the injured area immediately with water and seek medical advice immediately.
- Safety Data Sheets (SDS) are available on request.
- Use of this product should be limited to personnel trained in DNA amplification techniques.
- Workflow in the laboratory must be one-directional, beginning in the Extraction Area and moving to the Amplification and Detection Area. Do not return samples, equipment and reagents in the area where the previous step was performed.



Some components of this kit contain sodium azide as a preservative. Do not use metal tubing for reagent transfer.

## 6. SAMPLING AND HANDLING



Obtaining samples of biological materials for PCR-analysis, transportation, and storage are described in the manufacturer's handbook [1]. It is recommended that this handbook is read before starting work.

**AmpliSens<sup>®</sup> *C.trachomatis* / *Ureaplasma* / *M.genitalium*-MULTIPRIME-FRT** PCR kit is intended for analysis of DNA extracted with DNA extraction kits from the clinical material (urogenital, rectal and pharyngeal swabs; conjunctival discharge; prostate gland secretion; and urine samples).

## 7. WORKING CONDITIONS

**AmpliSens<sup>®</sup> *C.trachomatis* / *Ureaplasma* / *M.genitalium*-MULTIPRIME-FRT** PCR kit should be used at 18–25 °C.

## 8. PROTOCOL

### 8.1. DNA Extraction

It is recommended to use the following nucleic acid extraction kits:

- DNA-sorb-AM, **REF** K1-12-100-CE.
- For other nucleic acid extraction kits see Guidelines [2].

The DNA extraction of each test sample is carried out in the presence of **Internal Control-FL (IC)**.



Extract the DNA according to the manufacturer's protocol.

## 8.2. Preparing PCR

### 8.2.1. Preparing tubes for PCR

#### Variant FRT

The total reaction volume is **30 µl**, the volume of the DNA sample is **10 µl**.

1. Prepare the required number of the tubes with **PCR-mix-1-FL *C.trachomatis* / *Ureaplasma* / *M.genitalium*** and wax for amplification of DNA from clinical and control samples.
2. Add **10 µl** of **PCR-mix-2-FL-red** to the surface of the wax layer of each tube ensuring that it does not fall under the wax and mix with **PCR-mix-1-FL *C.trachomatis* / *Ureaplasma* / *M.genitalium***.

#### Variant FRT-100 F

The total reaction volume is **25 µl**, the volume of the DNA sample is **10 µl**.

1. Thaw the tube with **PCR-mix-2-FRT**. Mix the tubes with **PCR-mix-1-FL *C.trachomatis* / *Ureaplasma* / *M.genitalium*, PCR-mix-2-FRT, polymerase (TaqF)**. Sediment the drops by short centrifugation (1-2 s) using vortex.

Prepare the required number of the tubes for amplification of DNA from clinical and control samples.

2. For N reactions (including 2 controls), add to a new tube:

**10\*(N+1) µl of PCR-mix-1-FL *C.trachomatis* / *Ureaplasma* / *M.genitalium*,**

**5.0\*(N+1) µl of PCR-mix-2-FRT**

**0.5\*(N+1) µl of polymerase (TaqF).**

Vortex the tube, then centrifuge briefly. Transfer **15 µl** of the prepared mix into each tube.

Steps 3 and 4 are required in both variants.

3. Add **10 µl** of **DNA samples** obtained at the DNA extraction stage into prepared tubes.
4. Carry out the control amplification reactions:

**NCA** – Add **10 µl** of **DNA-buffer** to the tube labeled NCA (Negative Control of Amplification).

- C+** – Add **10 µl** of **Positive Control complex (C+)** to the tube labeled C+ (Positive Control of Amplification).
- C–** – Add **10 µl** of **the sample extracted from the Negative Control reagent** to the tube labeled C– (Negative control of Extraction).

### 8.2.2. Amplification

1. Create a temperature profile on your instrument as follows:

Table 1

#### AmpliSens-1 amplification program

Step	Rotor type instruments <sup>1</sup>			Plate type instruments <sup>2</sup>		
	Temperature, °C	Time	Cycles	Temperature, °C	Time	Cycles
1	95	15 min	1	95	15 min	1
2	95	5 s	5	95	5 s	5
	60	20 s		60	20 s	
	72	15 s		72	15 s	
3	95	5 s	40	95	5 s	40
	60	20 s (fluorescence detection)		60	30 s (fluorescence detection)	
	72	15 s		72	15 s	

Fluorescent signal is detected in the channels for the FAM, JOE, ROX and Cy5 fluorophores (other channels are enabled if several tests are simultaneously carried out in a single run).

2. Adjust the fluorescence channel sensitivity according to the *Important Product Information Bulletin* and Guidelines [2].
3. Insert tubes into the reaction module of the device.
4. Run the amplification program with fluorescence detection.
5. Analyze results after the amplification program is completed.

## 9. DATA ANALYSIS

Analysis of results is performed by the software of the real-time PCR instrument used by measuring fluorescence signal accumulation in four channels:

- The signal of the *Chlamydia trachomatis* DNA amplification product is detected in the channel for the FAM fluorophore.
- The signal of the *Ureaplasma* spp. (*U.parvum* and *U.urealyticum*) DNA amplification product is detected in the channel for the JOE fluorophore.
- The signal of the *Mycoplasma genitalium* DNA amplification product is detected in the channel for the ROX fluorophore.

<sup>1</sup> For example, Rotor-Gene 3000, Rotor-Gene 6000, Rotor-Gene Q, or equivalent

<sup>2</sup> For example, iCycler, iQ5, Mx3000P, Mx3000, or equivalent.



- The signal of the IC DNA amplification product is detected in the channel for the Cy5 fluorophore.

Results are interpreted by the crossing (or not-crossing) the fluorescence curve with the threshold line set at the specific level that corresponds to the presence (or absence) of a *Ct* value of the DNA sample in the corresponding column of the results grid.

Principle of interpretation is the following:

- ***Chlamydia trachomatis*** DNA is **detected** if the *Ct* value is determined in the results grid in the channel for the FAM fluorophore. Moreover, the fluorescence curve of the sample should cross the threshold line in the area of typical exponential growth of fluorescence.
- ***Ureaplasma spp. (U.parvum and U.urealyticum)*** DNA is **detected** if the *Ct* value is determined in the results grid in the channel for the JOE fluorophore. Moreover, the fluorescence curve of the sample should cross the threshold line in the area of typical exponential growth of fluorescence.
- ***Mycoplasma genitalium*** DNA is **detected** if the *Ct* value is determined in the results grid in the channel for the ROX fluorophore. Moreover, the fluorescence curve of the sample should cross the threshold line in the area of typical exponential growth of fluorescence.
- ***Chlamydia trachomatis, Ureaplasma spp. and Mycoplasma genitalium*** DNA are **not detected** in a sample if the *Ct* value is not determined (absent) in the channels for FAM, JOE and ROX fluorophores (fluorescence curve does not cross the threshold line), whereas the *Ct* value determined in the channel for the Cy5 fluorophore is less than the boundary *Ct* value specified in the *Important Product Information Bulletin*.
- The result is **invalid** if the *Ct* value is not determined (absent) in the channel for Cy5 fluorophore and in the channels for the FAM, JOE and ROX fluorophores. In such case, the PCR analysis of corresponding sample should be repeated.



Boundary *Ct* values are specified in the *Important Product Information Bulletin* enclosed to the PCR kit. See also Guidelines [2].

**The result of the analysis is considered reliable only if the results obtained for Positive and Negative Controls of amplification as well as for the Negative Control of extraction are correct (see Table 2).**

## Results for controls

Control	Stage for control	Ct value in the channel for fluorophore	
		FAM, JOE, ROX	Cy5
C–	DNA extraction	Absent	<boundary value
NCA	PCR	Absent	Absent
C+	PCR	<boundary value	<boundary value

## 10. TROUBLESHOOTING

Results of analysis are not taken into account in the following cases:

1. If the *Ct* value determined for the Positive Control of Amplification (C+) in the channels for the FAM and/or JOE and/or ROX fluorophores is greater than the boundary *Ct* value or absent, the amplification should be repeated for all samples in which the *Ct* value was absent in the channels for the FAM and/or JOE and/or ROX fluorophores.
2. If the *Ct* value is determined for the Negative Control of Amplification (NCA) and/or Negative Control of Extraction (C–) in the channels for the FAM and/or JOE and/or ROX fluorophores, the PCR analysis should be repeated for all samples in which the *Ct* value was determined in the channels for the FAM and/or JOE and/or ROX fluorophores.

If you have any further questions or if you encounter problems, please contact our Authorized representative in the European Community.

## 11. TRANSPORTATION

**AmpliSens® *C.trachomatis* / *Ureaplasma* / *M.genitalium*-MULTIPRIME-FRT** PCR kit should be transported at 2–8 °C for no longer than 5 days.

## 12. STABILITY AND STORAGE

All components of the **AmpliSens® *C.trachomatis* / *Ureaplasma* / *M.genitalium*-MULTIPRIME-FRT** PCR kit are to be stored at 2–8 °C when not in use (except for Polymerase (TaqF) and PCR-mix-2-FRT included in PCR kit variant FRT-100 F). All components of the **AmpliSens® *C.trachomatis* / *Ureaplasma* / *M.genitalium*-MULTIPRIME-FRT** PCR kit are stable until the expiry date stated on the label. The shelf life of reagents before and after the first use is the same, unless otherwise stated.



Polymerase (TaqF) and PCR-mix-2-FRT are to be stored at the temperature from minus 24 to minus 16 °C.



PCR-mix-1-FL *C.trachomatis* / *Ureaplasma* / *M.genitalium* is to be kept away from light.

## 13. SPECIFICATIONS

### 13.1. Sensitivity

Clinical material	Nucleic acid extraction kit	PCR kit	Microorganism	Sensitivity, GE/ml <sup>3</sup>
Urogenital swabs <sup>4</sup>	DNA-sorb-AM	PCR kit variants FRT and FRT-100 F	<i>Chlamydia trachomatis</i>	5x10 <sup>2</sup>
			<i>Ureaplasma</i> spp.	10 <sup>3</sup>
			<i>Mycoplasma genitalium</i>	10 <sup>3</sup>
Urine <sup>5</sup>	DNA-sorb-AM	PCR kit variants FRT and FRT-100 F	<i>Chlamydia trachomatis</i>	10 <sup>3</sup>
			<i>Ureaplasma</i> spp.	2x10 <sup>3</sup>
			<i>Mycoplasma genitalium</i>	2x10 <sup>3</sup>



The analytical sensitivity of each microorganism does not change even if two other microorganisms are present at high concentrations.

### 13.2. Specificity

The analytical specificity of the **AmpliSens<sup>®</sup> C.trachomatis / Ureaplasma / M.genitalium-MULTIPRIME-FRT** PCR kit is ensured by the selection of specific primers and probes as well as stringent reaction conditions. The primers and probes were checked for possible homologies to all sequences published in gene banks by sequence comparison analysis.

Nonspecific responses were absent while testing human DNA samples and DNA samples of microorganisms: *Gardnerella vaginalis*; *Lactobacillus* spp.; *Escherichia coli*; *Staphylococcus* spp.; *Streptococcus* spp.; *Candida albicans*; *Mycoplasma hominis*; *Ureaplasma urealyticum*; *Ureaplasma parvum*; *Mycoplasma genitalium*; *Chlamydia trachomatis*; *Neisseria* spp.; *Neisseria gonorrhoeae*; *Trichomonas vaginalis*; *Treponema pallidum*; *Toxoplasma gondii*; HSV types 1 and 2; CMV; HPV.

The clinical specificity of the **AmpliSens<sup>®</sup> C.trachomatis / Ureaplasma / M.genitalium-MULTIPRIME-FRT** PCR kit was confirmed in laboratory clinical trials.

## 14. REFERENCES

- Handbook "Sampling, Transportation, and Storage of Clinical Material for PCR diagnostics", developed by Federal Budget Institute of Science "Central Research

<sup>3</sup> The quantity of genome equivalents of microorganism per 1 ml of the clinical sample placed into the transport medium.

<sup>4</sup> Cervical and urethral swabs are to be placed into Transport Medium for Swabs (**REF** 956-CE, 987-CE) or Transport Medium with Mucolytic Agent (**REF** 952-CE, 953-CE).

<sup>5</sup> Pretreatment is required.














Institute for Epidemiology” of Federal Service for Surveillance on Consumers’ Rights Protection and Human Well-Being, Moscow, 2010.

2. Guidelines “Real-Time PCR Detection of STIs and Other Reproductive Tract Infections”, developed by Federal Budget Institute of Science “Central Research Institute for Epidemiology”.

## **15. QUALITY CONTROL**

In compliance with Federal Budget Institute of Science “Central Research Institute for Epidemiology” ISO 13485-Certified Quality Management System, each lot of the **AmpliSens® C.trachomatis / Ureaplasma / M.genitalium-MULTIPRIME-FRT** PCR kit has been tested against predetermined specifications to ensure consistent product quality.

## 16. KEY TO SYMBOLS USED

	Catalogue number		Sufficient for
	Batch code		Expiration Date
	<i>In vitro</i> diagnostic medical device		Consult instructions for use
	Version		Keep away from sunlight
	Temperature limitation	<b>NCA</b>	Negative control of amplification
	Manufacturer	<b>C-</b>	Negative control of extraction
	Date of manufacture	<b>C+</b>	Positive control of amplification
	Authorised representative in the European Community	<b>IC</b>	Internal control
	Caution		

### List of Changes Made in the Instruction Manual

VER	Location of changes	Essence of changes
23.06.11 RT	Cover page, text	The name of Institute was changed to Federal Budget Institute of Science “Central Research Institute for Epidemiology”
24.09.15 PM	Text	Corrections according to the template
	1. Intended use	The clinical material was specified
	6. Sampling and handling	
	3. Content, Footer	<b>REF</b> R-B46(iQ)-CE was deleted
	9. Data analysis	The sections were rewritten
10. Troubleshooting		