



For Professional Use Only

eSens *Ureaplasma parvum*/*urealyticum* QL PCR kit

REF ES3040A

Instructions for Use

1 INTENDED USE

eSens *Ureaplasma parvum*/*urealyticum* QL PCR kit is an *in vitro* nucleic acid amplification test for qualitative detection and differentiation of *Ureaplasma parvum* and *Ureaplasma urealyticum* DNA in the biological material (urogenital, rectal, and oropharyngeal swabs; conjunctival discharge; urine samples, prostate gland secretion) using real-time hybridization-fluorescence detection of amplified products.

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

2 PRINCIPLE OF PCR DETECTION

U.parvum / *U.urealyticum* detection by the polymerase chain reaction (PCR) is based on the amplification of the pathogen genome specific region using specific *U.parvum* / *U.urealyticum* 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.

eSens *Ureaplasma parvum*/*urealyticum* QL 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.

eSens *Ureaplasma parvum*/*urealyticum* QL PCR kit uses "hot-start," which greatly reduces the frequency of nonspecifically primed reactions. "Hot-start" is guaranteed by using chemically modified polymerase (TaqF). The chemically modified polymerase (TaqF) is activated by heating at 95 °C for 15 min.

The PCR kit contains the system for prevention of contamination by amplicons using the enzyme uracil-DNA-glycosylase (UDG) and dUTP. The enzyme UDG recognizes and catalyzes the destruction of the DNA containing deoxyuridine, but has no effect on DNA containing deoxythymidine. Deoxyuridine

is absent in the authentic DNA, but is always present in amplicons, because dUTP is a part of dNTP mixture in the reagents for the amplification. Due to the deoxyuridine containing contaminating amplicons are sensitive to the destruction by UDG before the DNA-target amplification. So the amplicons cannot be amplified.

The enzyme UDG is thermolabile. It is inactivated by heating at temperature above 50 °C. Therefore, UDG does not destroy the target amplicons which are accumulated during PCR.

The results of amplification are registered in the following fluorescence channels:

Table 1

Channel for fluorophore	FAM	JOE	ROX
DNA-target	<i>U. parvum</i> DNA	<i>U. urealyticum</i> DNA	Internal Control-FL (IC) DNA
Target gene	Urease gene	Urease gene	Artificially synthesized sequence

3 CONTENT

eSens *Ureaplasma parvum/urealyticum* QL PCR kit (ES3040A) contains:

Reagent	Description	Volume, ml	Quantity
PCR-mix-1-FL <i>U.parvum</i> / <i>U.urealyticum</i>	clear liquid from colorless to light lilac colour	1.2	1 tube
PCR-mix-2-FRT	colorless clear liquid	0.3	2 tubes
Polymerase (TaqF)	colorless clear liquid	0.03	2 tubes
Positive Control complex (C+)	colorless clear liquid	0.2	1 tube
DNA-buffer	colorless clear liquid	0.5	1 tube
Negative Control (C-)*	colorless clear liquid	1.2	1 tube
Internal Control-FL (IC)**	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.

eSens *Ureaplasma parvum/urealyticum* QL PCR kit is intended for 110 reactions (including controls).

4 ADDITIONAL REQUIREMENTS

- DNA extraction kit.
- Transport medium.
- Disposable powder-free gloves and a laboratory coat.
- Pipettes (adjustable).

- Sterile pipette tips with aerosol barriers (up to 100 µl).
- Tube racks.
- Vortex mixer.
- Desktop centrifuge with rotor for 2-ml reaction tubes.
- PCR box.
- Real-time instruments (for example, Rotor-Gene Q (QIAGEN, Germany), CFX 96 Touch, CFX 96 Opus (Bio-Rad, USA), QuantStudio 5 (Thermo Fisher Scientific), or equivalent).
- Disposable polypropylene PCR tubes (0.1 or 0.2 ml)
 - 0.2-ml PCR tubes with optical transparent domed or flat caps if a plate-type instrument is used;
 - 0.2-ml 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 at the temperature from minus 24 to minus 16 °C.

5 GENERAL PRECAUTIONS

The user should always pay attention to the following:

- Use sterile pipette tips with aerosol filters 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 inhalation of vapors, samples and reagents contact with the skin, eyes, and mucous membranes. Harmful if swallowed. If these solutions come into contact, rinse the injured area immediately with water and seek medical advice if necessary.
- 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

eSens Ureaplasma parvum/urealyticum QL PCR kit is intended for analysis of the DNA extracted with DNA extraction kits from the clinical material (urogenital, rectal, and oropharyngeal swabs, conjunctival

discharge, urine samples (a sediment of the first portion of the morning specimen), prostate gland secretion).

7 WORKING CONDITIONS

eSens Ureaplasma parvum/urealyticum QL PCR kit should be used at 18–25 °C.

8 PROTOCOL

8.1 DNA extraction

Any commercial nucleic acid extraction kit, if IVD-CE validated for the indicated specimen types, could be used.

Ecoli Dx, s.r.o. recommends:

- For the manual extraction

- **DNA-sorb-AM** (K1-12-100-CE)

- For the automatic extraction

- **ePure STD DNA Extraction Kit (E2007)**

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

Please carry out nucleic acid extraction according to the manufacture´s instruction.

8.2 Preparing PCR

8.2.1 Preparing tubes for PCR

The type of tubes depends on the PCR instrument used for analysis. Use disposable filter tips for adding reagents, DNA and control samples into tubes.

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

1. Thaw the tube with PCR-mix-2-FRT. Vortex the tubes with PCR-mix-1-FL *U.parvum* / *U.urealyticum*, PCR-mix-2-FRT, and polymerase (TaqF) then centrifuge briefly. Take the required number of the tubes/strips for amplification of DNA obtained from clinical and control samples.
2. For N reactions (including 2 controls of amplification) add to a new tube:
10·(N+1) µl of PCR-mix-1-FL *U.parvum* / *U.urealyticum*,
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 mixture into each tube.**
3. Add **10 µl of DNA samples** obtained at the DNA extraction stage.
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 Negative Control (C–) 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 2

eSens-1 amplification program

	Rotor-type Instruments (e.g Rotor-Gene Q or equivalent)			Plate-type Instruments (e.g CFX 96 Touch, CFX 96 Opus, QuantStudio 5 or equivalent.)		
Step	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 <i>Fluorescence acquiring</i>		60	30 s <i>Fluorescence acquiring</i>	
	72	15 s		72	15 s	

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

2. Adjust the fluorescence channel sensitivity.
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.

8.3 Instrument Settings

Test settings for rotor-type instruments

Channel	Calibrate/Gain Optimisation	Threshold	Dynamic tube	Slope Correct	More Settings/ Outlier Removal
FAM/Green	from 5 FI to 10 FI	0.1	On	Off	0%
JOE/Yellow	from 4 FI to 8 FI	0.1	On	Off	5%
ROX/Orange	from 4 FI to 8 FI	0.1	On	Off	5%

Test settings for plate-type instruments

Set the heating/cooling **Ramp Rate 2,5 °C/s**.

Channel	Threshold
FAM JOE/HEX	For each channel in Log Scale set the threshold line at the level of 10-20 % of maximum fluorescence obtained for the Positive Control of Amplification (C+) in the last amplification cycle

9 DATA ANALYSIS

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

- The signal of the *U.parvum* DNA amplification product is detected in the channel for the FAM fluorophore.
- The signal of the *U.urealyticum* DNA amplification product is detected in the channel for the JOE fluorophore.
- The signal of the IC DNA amplification product is detected in the channel for the ROX 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 *Ct* value of the DNA sample in the corresponding column of the result grid.

Principle of interpretation is the following:

- *Ureaplasma parvum* DNA is **detected** in a sample if the *Ct* value is determined in the result 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 urealyticum* DNA is **detected** in a sample if the *Ct* value is determined in the result 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.
- *Ureaplasma parvum* and *Ureaplasma urealyticum* DNA are **not detected** in a sample if the *Ct* value is not determined (absent) in the result grid in the channels for the FAM and JOE fluorophores (the fluorescence curve does not cross the threshold line), whereas the *Ct* value in the channel for ROX fluorophore is less than the specified boundary *Ct* value.
- The result is invalid if *Ct* value is not determined (absent) in the channels for FAM, JOE and ROX fluorophores. In such cases, the PCR analysis should be repeated for such samples.

The result of the analysis is considered reliable only if the results obtained for the Positive and Negative Controls of amplification as well as for the Negative Control of extraction are correct (see Table 3 and 4).

Table 3

Results for controls

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

Table 4

Boundary Ct values

Sample	Rotor-type instrument			Plate-type instrument		
	Channel for fluorophore					
	FAM	JOE	ROX	FAM	JOE	ROX
C+	33	33	33	36	36	36
C-	Ct is absent		33	Ct is absent		36
NCA	Ct is absent			Ct is absent		
Test samples	-	-	33	-	-	36

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 fluorophores is greater than the boundary Ct value or absent, the amplification should be repeated for all samples in which the boundary Ct is absent in the channels for the FAM and/or JOE 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 fluorophores, the PCR analysis should be repeated for all samples in which the Ct value is determined in the channels for the FAM and/or JOE fluorophores.

11 TRANSPORTATION

eSens Ureaplasma parvum/urealyticum QL PCR kit should be transported at 2–8 °C for no longer than 10 days.

12 STABILITY AND STORAGE

All components of the **eSens Ureaplasma parvum/urealyticum QL PCR kit** (except for polymerase (TaqF) and PCR-mix-2-FRT) are to be stored at 2–8 °C when not in use.

All components of the **eSens Ureaplasma parvum/urealyticum QL PCR kit** are stable until labeled expiry date. PCR kit can be stored without unpacking at 2 to 8 °C for 3 months from the date of manufacture before opening. Once opened, PCR kit should be unpacked in accordance with the storage temperatures for each component. The shelf life of opened reagents is the same as that of unopened reagents, unless otherwise stated.

NOTE: Polymerase (TaqF) and PCR-mix-2-FRT are to be stored at temperature from minus 24 to minus 16 °C when not in use

NOTE: PCR-mix-1-FL *U.parvum/U.urealyticum* is to be kept away from light.

13 SPECIFICATIONS

13.1 Sensitivity

Clinical material	Nucleic acid extraction kit	Microorganism	Analytical sensitivity, GE/ml*
Urogenital swabs**	DNA-sorb-AM ePure STD DNA Extraction Kit	<i>Ureaplasma parvum</i>	10 ³
		<i>Ureaplasma urealyticum</i>	10 ³
Urine (pretreatment is required)	DNA-sorb-AM ePure STD DNA Extraction Kit	<i>Ureaplasma parvum</i>	5x10 ³
		<i>Ureaplasma urealyticum</i>	5x10 ³

* Genome equivalents (GE) of the pathogen agent per 1 ml of a sample placed in the transport medium.

** Urogenital swabs are to be placed into **Transport Medium for swabs** or **Transport Medium with Mucolytic Agent**.

13.2 Specificity

The analytical specificity of **eSens Ureaplasma parvum/urealyticum QL PCR kit** is ensured by selection of specific primers and probes as well as stringent reaction conditions. The primers and probes were checked for possible homologies to all sequences deposited in gene banks by sequence comparison analysis.

Nonspecific responses were absent while testing human DNA samples as well as a DNA panel of the following 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.*, *Trichomonas vaginalis*, *Neisseria gonorrhoeae*, *Treponema pallidum*, *Toxoplasma gondii*, *HSV 1 and 2*, *CMV*, and *HPV*.

The clinical specificity of **eSens Ureaplasma parvum/urealyticum QL PCR kit** was confirmed in laboratory clinical trials.

14 QUALITY CONTROL

The production process, including batch release, is carried out in accordance with an established quality management system certified according to ISO 13485.

15 KEY TO SYMBOLS USED

	Catalogue number		Caution
	Batch code		Contains sufficient for <n> tests
	<i>In vitro</i> diagnostic medical device		Use-by Date
	Version		Consult instructions for use
	Temperature limit		Keep away from sunlight
	Manufacturer	NCA	Negative control of amplification
	Date of manufacture	C-	Negative control of extraction
	Authorized representative in the European Community	C+	Positive control of amplification
		IC	Internal control

List of Changes Made in the Instruction Manual

VER	Location of changes	Essence of changes
01_04/2022		
02_02/2025	Intended Use Sensitivity	Modified terminology of biological / clinical material. Modification of urogenital swabs are to be placed into Transport Medium for swabs or Transport Medium with Mucolytic Agent.
03_10/2026	Transportation	Transportation period was changed from 5 to 10 days.

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