

DECLARATION OF VERIFICATION

Diagnostic medical devices *in vitro*

GeneProof

Manufacturer
GeneProof a.s.,
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GeneProof a.s hereby declares that following products are verified together with the devices:

GeneProof PCR kits	Bloodborne Infections																				
	croBEE Real-Time PCR System	Applied Biosystems 7300 Real-Time PCR System	Applied Biosystems 7500 Real-Time PCR System	AriaMx Real-Time PCR System	CFX Connect™ Real-Time PCR Detection System	CFX96™ / Dx Real-Time PCR Detection System	DT lite Real-Time PCR System	LightCycler® 2.0	LightCycler® 480	LineGene 9600	LineGene 9600 Plus	Mic qPCR Cycler	Montania 4896 Real-Time PCR termocykler	QuantStudio™ 3 Real-Time PCR System	QuantStudio™ 5 Real-Time PCR System	Rotor-Gene 3000	Rotor-Gene 6000	Rotor-Gene Q	SLAN® Real-Time PCR System	StepOne™ / StepOne Plus™ Real-Time PCR System	Stratagene Mx3005P qPCR System
GeneProof Hepatitis B Virus (HBV) PCR Kit																					X
GeneProof Hepatitis C Virus (HCV) Diagnostic PCR Kit							X	-		-			X	X	X				X	-	X
GeneProof HIV type 1 (HIV-1) PCR Kit							X	-		-			X			X				-	X
Immunocompromised / Transplanted																					
GeneProof Aspergillus PCR Kit		X			X		X	X					X	X				X		X	X
GeneProof BKJC Virus (BKJC) PCR Kit		X			X		X	X					X	X	X			X		X	X
GeneProof Cytomegalovirus (CMV) PCR Kit							X														X
GeneProof Epstein-Barr Virus (EBV) PCR Kit							X											X			X
GeneProof Herpes Simplex Virus 1 (HSV-1) PCR Kit							X						X								X
GeneProof Herpes Simplex Virus 2 (HSV-2) PCR Kit							X						X								X
GeneProof Herpes Simplex Virus (HSV-1/2) PCR Kit							X						X								X
GeneProof Human Herpes Virus 6/7 (HHV-6/7) PCR Kit		X			X		X	X					X	X			X	X		X	X
GeneProof Human Herpesvirus 8 (HHV-8) PCR Kit							X	-		-			X	X							X
GeneProof Parvovirus B19 PCR Kit							X						X								X
GeneProof Varicella-Zoster Virus (VZV) PCR Kit							X						X				X				X
Sexually Transmitted Infections																					
GeneProof Chlamydia trachomatis PCR Kit													X								X
GeneProof CT/NG/MG Multiplex PCR Kit		X			X		X	X		-			X	X			X		X		X
GeneProof Human Papillomavirus (HPV) Screening PCR Kit		-	1)	1)	2)		X	-	2)	-		X	X	6)	1)	-	-	7)	3)	-	1)
GeneProof MH/UV/UP Multiplex PCR Kit		X		-	X		X	X	X	-			X	X		X	X	X	X		X
GeneProof Mycoplasma genitalium/hominis PCR Kit		X	X		4)		X	X	X				X	4)			4)				X
GeneProof Neisseria gonorrhoeae PCR Kit							X						X								X
GeneProof Treponema pallidum PCR Kit				-			X	X		-			X				X				X
GeneProof Trichomonas vaginalis PCR Kit				-			X	-		-			X								X
GeneProof Ureaplasma PCR Kit		X	X		5)		X	X					X	5)			5)				X

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Legend:	
Verified	
Not compatible system (for technical reasons)	X
Not supported system (not verified)	X
Not tested yet	-

- 1) The device allows detection of high risk HPV with differentiation of HPV 16 and HPV 18
- 2) The device allows detection of high risk HPV without differentiation.
- 3) The device allows detection of high risk HPV with differentiation of HPV 16.
- 4) The device allows detection of *M. genitalium* only.
- 5) The device does not allow to distinguish between *U. parvum* and *U. urealyticum*.

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- 6) Only device model which allows 3-channel detection (FAM, HEX, Cy5).
- 7) The device allows detection of high-risk HPV and presence HPV18, 16 and 45 but it does not distinguish coinfection of types 16, 18 and 45 respect to the cross-talk between the channels TEX/Cy5 and Cy5/CY5.5
- 8) The device allows detection of high risk HPV with differentiation of HPV 18.

Products together with the devices are verified to comply with the requirements of the specified or expected use (i.e. it complies with the verification requirements), according to the Council and European Parliament Directive 98/79/EC dated October 27, 1998, on *in vitro* diagnostic medical devices, requirements of which were adopted in the Czech Government Regulation No. 56/2015 Coll. establishing technical requirements for *in vitro* diagnostic medical devices.

A2.12.2023



Mgr. Petra ŠVÁSTOVÁ
QA/QC Department
QC Manager
(Name, position and signature of authorized person)

Manufacturer's stamp:

