

## Measles and Rubella National Laboratory or sub-National Laboratory Checklist for WHO Accreditation Section 1: General Review & Overall Findings

Dates of 02	1/01/2024 -	On-site Oesk R	eview	Accreditation for	r		
Review: 32	1/12/2024			calendar year	: 2026		
Name of L	Laboratory	of Communicable I	Diseases				
Laboratory: Health Board / Terviseamet							
Address: Paldiski mnt 81, 10614 Tallinn							
				Country ESTONIA			
Phone: 372794	<b>43500</b> E1	mail: <u>oksana.joamet</u>	s@terviseamet.ee	Website: www.tervisean	<u>iet.ee</u>		
		<u>liisi.kink@terv</u>	riseamet.ee				
Head of Institute	: Bı	rigit Lao	Head of Departm	nent: Liisi Kink			
Head of Measles	and						
Rubella Laborato	ory: Ol	ksana Joamets					
Technical Supervisor: Oksana Joamets							
Number of laboratories under supervision: 1							
Name of							
Reviewer(s): Dr. Myriam Ben Mamou & Dovile Videbaek, WHO/Europe							
Name of National Accreditation Authority (if appropriate) and current accreditation status from this							

Name of National Accreditation Authority (if appropriate) and current accreditation status from this authority (e.g. ISO15189 or other, provide documentation):

Estonian Accreditation Centre, EVS-EN ISO 15189:2022.

Accreditation Certificate No M004. Accreditation is valid until 29.06.2029

(http://eak.ee/dokumendid/pdf/kasitlusala/M004.pdf)

# **Summary of Accreditation Review**

#### Recommendations (check one):

$\boxtimes$	Accredit: Laboratory meets all criteria					
	Measles:	Rubella:  Serology  Detection RT-PCR/RT-qPCR  Sequencing Cell Culture Rubella testing not done				
	<b>Provisionally accredit</b> : Laboratory passed the most recent proficiency test, but needs to strengthen one or more of the remaining criteria (check off relevant criteria):					
	Measles:  Serology Detection RT-PCR/RT-qPCR Sequencing Cell Culture Measles testing not done  Do not accredit: Laboratory did not pasother performance indicators.	Rubella: Serology Detection RT-PCR/RT-qPCR Sequencing Cell Culture Rubella testing not done ss the most recent proficiency test and/or did not meet				
	Measles:  Serology  Detection RT-PCR/RT-qPCR  Sequencing Cell Culture Measles testing not done	Rubella: Serology Detection RT-PCR/RT-qPCR Sequencing Cell Culture Rubella testing not done				

#### **Main General Findings:**

### **Findings:**

1.	Confirmatory test results of Sub-national Laboratories' referred swithin 14 days for $\geq 80\%$ of specimens received:	100%	
2.	The laboratory reports monthly to WHO ( $\geq$ 80%):	Completeness: Timeliness:	100%
3.	IgM tests are performed on at least 50 specimens annually:	Measles: Rubella:	59 248
4a.	Measles and Rubella IgM test results conducted for primary diagnostics are reported by the laboratory within 4 days of receipt, for $\geq 80\%$ of specimens	Measles: Rubella:	95% 100%
4b.	If molecular virus detection by real-time RT-PCR is performed in addition to serology, positive test results are reported by the laboratory within 4 days of receipt, for ≥80% of specimens	Measles: Rubella:	100% NA
5.	The accuracy of IgM detection is $\geq$ 90% (NLs sending sera to RRL and SNLs to NLs for confirmatory testing) - <b>50 measles and 4 rubella samples</b>	Measles: Rubella:	98% 100%
6a.	At least one run per month is performed for measles and rubella detection Realtime RT-PCR	Measles: Rubella:	Yes NA
6b.	At least one run per quarter is performed for measles and rubella GT RT-PCR & sequencing	Measles: Rubella:	Yes NA
7.	Score on most recent WHO Serology proficiency test is $\geq 90\%$ : PT panel number: <b>02401</b> Date reported: <b>20/03/2025</b>	Measles:	Second test 100%
	Parformance on most recent WHO Malacular FOA	Rubella:  Measles Mol detection:	Second test 100% Passed
	Performance on most recent WHO Molecular EQA is passed ( <b>P</b> ) /re-test ( <b>R</b> ) (provisionally passed)/fail ( <b>F</b> )	Measles Sequencing:	Passed
8.	Panel number for measles: 24-1 Date reported: 29/11/2024	Rubella Mol detection:	NA
	Panel number for rubella: Date reported: dd/mm/yyyy	Rubella Sequencing:	NA
9.	For labs that conduct virus sequencing: Virus sequencing is compand AND reported to WHO through MeaNS or RubeNS within 2 mor receipt of specimens, for ≥80% of the specimens appropriate for ganalysis	oleted nths of MeaNS;	100% (2/2) NA
10.	Score from on-site review of laboratory operating procedures and		

#### **Conclusion**

- Good performance for serology; excellent for measles rRT-PCR and sequencing
- Excellent monthly data reporting to WHO.