Date of report: 23/04/2025

JITM Diagnostics

Patient data				
Name	MRS. ARTI	Patient ID		
Birthday	11/01/1993 S			2504047152/NOD
Age at sample date	32.3		•	23/04/2025
Gestational age	12 + 6			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 89.2	diabetes	no pregnancies		
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Corr. MoM	rr. MoM Gestational age 12 + 5		
PAPP-A 2.5 mIU/m	I 0.89	Method CRL Robinson		
fb-hCG 29.1 ng/ml	0.75	Scan date 22/04/2025		
Risks at sampling date			Crown rump length in mm 66	
Age risk	1:468	Nuchal translucency MoM 0.89		
Biochemical T21 risk	1:4129	Nasal bone present		
Combined trisomy 21 risk	risk <1:10000		Sonographer	
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT		MD
Risk 1:10		Trisomy 21	ated risk for Trisomy 21	
1:1000 1:10000 1:10000 1:110000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.		translucency) is below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approaches and have no diagnostic value! The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)). The laboratory can not be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic value!		

Sign of Physician