Prisca 5.2.0.13

Date of report: 9/04/2025

## JITM Diagnostics

Patient data						
Name	MRS. RUMA			Patient ID		
Birthday	20/09/1991			Sample ID		2504016827/NOD
Age at sample date	e 33.5			Sample Date	9	8/04/2025
Gestational age 12 + 2						
Correction factors						
Fetuses	1	IVF		no	Previous trisomy 21	no
Weight	73	diabetes		no	pregnancies	
Smoker	no	Origin		Asian		
Biochemical data				Ultrasound data		
Parameter	Value	C	orr. MoM	Gestational	age	12 + 2
PAPP-A	2.1 mIU/m	I	0.74	Method		CRL Robinson
fb-hCG	20.1 ng/ml		0.46			8/04/2025
Risks at sampling date			Crown rump length in mm 59			
Age risk	1:363		Nuchal translucency MoM		1.43	
Biochemical T21 risk	1.0011			Nasal bone		present
Combined trisomy 21 risk 1:6044			Sonographer DR. ANKIT KHANDELWAL MBBS DNB			
Trisomy 13/18 + NT 1:6875			Qualifications in measuring NT MD			
Risk 1:10 1:250 Cut off 1:100 1:250 Cut off 1:1000 1:100 1:10000 1:10000 1:10000 1:100			Trisomy 21 The calculated risk for Trisomy 21 (with nuchal 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 6044 women with the same data, there is one woman with a trisomy 21 pregnancy and 6043 women with not affected pregnancies. 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!			
Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with nuchal						

The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:6875, which represents a low risk.

Sign of Physician