Prisca 5.2.0.13

Date of report: 15/02/2025

Patient data					
Name	MRS. MOHINI TRIPATHI				
Birthday	2/04/1995		Sample ID	2502026971/NOD	
Age at sample date 29.9		Sample Date 14/02/2025			
Gestational age 12 + 4					
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 no	
Weight	56	diabetes			
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter V	Value Corr. MoM		Gestational age 12 + 3		
PAPP-A 3.46	mIU/m	0.78	Method CRL Robinson		
	ng/ml	1.49	Scan date 13/02/2025		
Risks at sampling date			Crown rump length in mm 61		
Age risk	1:657		Nuchal translucency MoM 0.95		
Biochemical T21 risk			Nasal bone present		
Combined trisomy 21 risk 1:4370			Sonographer		
Trisomy 13/18 + NT				Qualifications in measuring NT	
KISK 1:10			Trisomy 21	ated risk for Trisomy 21 (with nuchal	
1:1000 1:250 Cut off 1:10000 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 below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 4370 women with the same data, there is one woman with a trisomy 21 pregnancy and 4369 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!		
translucency) is < 1:10000, which represents a low risk.					