Prisca 5.2.0.13

Date of report: 1/02/2025

Patient data				
Name	MRS. SHABANA			
Birthday	4/06/1990	Sample ID		2501051457/NOD
Age at sample date	34.7)	31/01/2025
Gestational age	12 + 2			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 76	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	nta	
Parameter Value	Corr. MoM	Gestational age 11 + 4		
PAPP-A 3.45 mIU/m	nl 1.28	Method CRL Robinson		
fb-hCG 36.5 ng/ml	0.85			
Risks at sampling date			length in mm	50.4
Age risk	1:290	Nuchal translucency MoM 1.03		
Biochemical T21 risk	1:4264	Nasal bone		present
Combined trisomy 21 risk	,		Sonographer	
isomy 13/18 + NT <1:10000		Qualifications in measuring NT Trisomy 21		
1:10 1:250 Cut off 1:1000 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.		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 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