

Prisca

5.1.0.17

Date of report: 04-09-2025

NA

Patient data			
Name	W/O ROHIT		Patient ID
Birth day	12-06-1995		Sample ID
Age at sample date	30.2		Sample Date
Gestational age	11 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	74	diabetes	no
Smoker	no	Origin	Asian
			Previous trisomy 21 pregnancies
			unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.36 mIU/mL	0.61	11 + 4
fb-hCG	44.65 ng/ml	0.98	Method
Risks at sampling date			CRL Robinson
Age risk	1:605		Scan date
Biochemical T21 risk	1:1237		02-09-2025
Combined trisomy 21 risk	1:6983		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		50.1
			Nuchal translucency MoM
			0.52
			Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			MD
Risk		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 6983 women with the same data, there is one woman with a trisomy 21 pregnancy and 6982 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 translucency) is < 1:10000, which represents a low risk.			

Sign of Physician

below cut off
 Below Cut Off, but above Age Risk
 above cut off