

Prisca 5.1.0.17
Date of report: 29/09/25

NA

Patient data			
Name	Ms. TARALI PATAR		Patient ID 0692509270107
Birthday	22/02/96		Sample ID A1738430
Age at sample date	29.6		Sample Date 27/09/25
Gestational age	13 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	58.7	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 13 + 5
PAPP-A	4.13 mIU/mL	0.63	Method CRL Robinson
fb-hCG	29.88 ng/mL	1.02	Scan date 26/09/25
Risks at sampling date			Crown rump length in mm 80.04
Age risk	1:709		Nuchal translucency MoM 2.02
Biochemical T21 risk	1:1420		Nasal bone present
Combined trisomy 21 risk	1:104		Sonographer NA
Trisomy 13/18 + NT	1:1538		Qualifications in measuring NT NA
Trisomy 21			
The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk. After the result of the Trisomy 21 test (with NT) it is expected that among 104 women with the same data, there is one woman with a trisomy 21 pregnancy and 103 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 1:1538, which represents a low risk.			

Sign of Physician

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