

Prisca 5.1.0.17
Date of report: 01/10/24

Patient data			
Name	Mrs. SANJIVANI TATHE	Patient ID	0672409300163
Birthday	26/03/04	Sample ID	A1189070
Age at sample date	20.5	Sample Date	30/09/24
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	49	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	7.92 mIU/mL	1.39	Gestational age 12 + 4
fb-hCG	121.02 ng/mL	2.92	Method CRL Robinson
			Scan date 28/09/24
Risks at sampling date			Crown rump length in mm 63
Age risk		1:1094	Nuchal translucency MoM 0.74
Biochemical T21 risk		1:920	Nasal bone present
Combined trisomy 21 risk		1:4887	Sonographer NA
Trisomy 13/18 + NT		<1:10000	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 4887 women with the same data, there is one woman with a trisomy 21 pregnancy and 4886 women with not affected pregnancies. The free beta HCG level is high. 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