

Prisca 5.1.0.17
Date of report: 11-08-2025

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Patient data			
Name	Mrs. MANINI NAYAK.		Patient ID
Birthday	27-10-1989	Sample ID	
Age at sample date	35.8	Sample Date	
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	70	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.97 mIU/mL	0.86	
fb-hCG	33.8 ng/mL	1.07	
Risks at sampling date			
Age risk	1:236		Gestational age
Biochemical T21 risk	1:898		Method
Combined trisomy 21 risk	1:4389		Scan date
Trisomy 13/18 + NT	<1:10000		Crown rump length in mm
			Nuchal translucency MoM
			Nasal bone
			Sonographer
			Qualifications in measuring NT
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 4389 women with the same data, there is one woman with a trisomy 21 pregnancy and 4388 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