

Prisca 5.1.0.17
Date of report: 11-03-2025

DR
S MANJULA MBBS

Patient data			
Name Mrs. PRIYANKA SURYAWANSHI		Patient ID 0372503100066	
Birthday 28-09-1991		Sample ID A1501025	
Age at sample date 33.4		Sample Date 10-03-2025	
Gestational age 13 + 5			
Correction factors			
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 72	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 13 + 1
PAPP-A	4.98 mIU/mL	1.01	Method CRL Robinson
fb-hCG	50.12 ng/mL	1.74	Scan date 06-03-2025
Risks at sampling date			Crown rump length in mm 71
Age risk 1:390			Nuchal translucency MoM 0.62
Biochemical T21 risk 1:669			Nasal bone unknown
Combined trisomy 21 risk 1:3588			Sonographer DR S MANJULA MBBS
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT DMCH
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 3588 women with the same data, there is one woman with a trisomy 21 pregnancy and 3587 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