

Prisca 5.2.0.13

Date of report: 2/07/2025

Patient data			
Name		MRS. SWETA	Patient ID
Birthday		22/05/1988	Sample ID
Age at sample date		37.1	2507002552/NOD
Gestational age		13 + 3	Sample Date
			2/07/2025
Correction factors			
Fetuses	1	IVF	no
Weight	58	diabetes	no
Smoker	no	Origin	Asian
			Previous trisomy 21 pregnancies
			no
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	10.1 mIU/ml	1.69	13 + 2
fb-hCG	56.1 ng/ml	1.37	Method
Risks at sampling date			CRL Robinson
Age risk	1:172		Scan date
Biochemical T21 risk	1:1421		1/07/2025
Combined trisomy 21 risk	1:6495		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		74.7
			Nuchal translucency MoM
			0.71
			Nasal bone
			present
			Sonographer
			DR. NEERJA CHOPRA
			Qualifications in measuring NT
			M.D
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 6495 women with the same data, there is one woman with a trisomy 21 pregnancy and 6494 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.			

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