

Prisca 5.1.0.17
Date of report: 30-11-2025

Patient data			
Name	Mrs. P SARITHA		Patient ID
Birthday	08-08-1997		Sample ID
Age at sample date	28.3		Sample Date
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	65	diabetes	no
Smoker	no	Origin	Asian
			Previous trisomy 21 pregnancies
			unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.92 mIU/mL	0.38	13 + 2
fb-hCG	31.8 ng/mL	0.99	Method
			CRL Robinson
			Scan date
			27-11-2025
Risks at sampling date			Trisomy 21
Age risk			1:798
Biochemical T21 risk			1:449
Combined trisomy 21 risk			1:2404
Trisomy 13/18 + NT			<1:10000
			Crown rump length in mm
			73
			Nuchal translucency MoM
			0.94
			Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			Sonographer
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 2404 women with the same data, there is one woman with a trisomy 21 pregnancy and 2403 women with not affected pregnancies. The PAPP-A level is low. 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