

Prisca 5.1.0.17
Date of report: 18-03-2025

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Patient data			
Name	Mrs. G.SARITHA		Patient ID
Birthday	06-05-1998	Sample ID	0842503150011
Age at sample date	26.9	Sample Date	A2172244
Gestational age	12 + 3		15-03-2025
Correction factors			
Fetuses	1	IVF	no
Weight	57	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.19 mIU/mL	0.79	12 + 3
fb-hCG	58.11 ng/mL	1.34	Method
			CRL Robinson
			Scan date
			15-03-2025
Risks at sampling date			Trisomy 21
Age risk	1:865		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
Biochemical T21 risk	1:1629		After the result of the Trisomy 21 test (with NT) it is expected that among 8979 women with the same data, there is one woman with a trisomy 21 pregnancy and 8978 women with not affected pregnancies.
Combined trisomy 21 risk	1:8979		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!
Trisomy 13/18 + NT	<1:10000		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