

Prisca 5.1.0.17
Date of report: 21-08-2025

PRAGATHI

Patient data			
Name	Mrs. VARA LAXMI	Patient ID	0012508190504
Birthday	08-04-1988	Sample ID	B3535511
Age at sample date	37.4	Sample Date	19-08-2025
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	49	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.01 mIU/mL	0.56	12 + 5
fb-hCG	39.14 ng/mL	0.91	Method
			CRL Robinson
			Scan date
			19-08-2025
Risks at sampling date			Crown rump length in mm
Age risk	1:157		65
Biochemical T21 risk	1:300		Nuchal translucency MoM
Combined trisomy 21 risk	1:1715		0.60
Trisomy 13/18 + NT	<1:10000		Nasal bone
			present
			Sonographer
			N A
			Qualifications in measuring NT
			MD
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 1715 women with the same data, there is one woman with a trisomy 21 pregnancy and 1714 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

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