

Prisca 5.1.0.17
Date of report: 28-11-2025

Patient data			
Name Mrs. LAVANYA W/O PRASHANTH		Patient ID 0012511270362	
Birthday 15-03-2001		Sample ID A2152861	
Age at sample date 24.7		Sample Date 27-11-2025	
Gestational age 13 + 0			
Correction factors			
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 71	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 13 + 0
PAPP-A	0.96 mIU/mL	0.25	Method CRL Robinson
fb-hCG	38.74 ng/mL	1.10	Scan date 27-11-2025
Risks at sampling date			Crown rump length in mm 70
Age risk 1:989			Nuchal translucency MoM 0.80
Biochemical T21 risk 1:118			Nasal bone present
Combined trisomy 21 risk 1:862			Sonographer NA
Trisomy 13/18 + NT 1:7461			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 862 women with the same data, there is one woman with a trisomy 21 pregnancy and 861 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 1:7461, which represents a low risk.			

Sign of Physician