

Prisca 5.1.0.17
Date of report: 11-12-2025

Patient data			
Name Mrs. K SUNITHA W/O SANTHOSH CH		Patient ID 0012512100327	
Birthday 12-07-1991		Sample ID A2152986	
Age at sample date 34.4		Sample Date 10-12-2025	
Gestational age 12 + 0			
Correction factors			
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 50	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 11 + 5
PAPP-A	1.95 mIU/mL	0.50	Method CRL Robinson
fb-hCG	44.2 ng/mL	0.88	Scan date 08-12-2025
Risks at sampling date			Crown rump length in mm 52
Age risk 1:302			Nuchal translucency MoM 0.72
Biochemical T21 risk 1:460			Nasal bone present
Combined trisomy 21 risk 1:2697			Sonographer NA
Trisomy 13/18 + NT <1:10000			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 2697 women with the same data, there is one woman with a trisomy 21 pregnancy and 2696 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 held 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