

Prisca 5.1.0.17
Date of report: 21-10-2025

N A

Patient data			
Name Mrs. B SAVITHA W/O SANTHOSH		Patient ID 0012510210148	
Birthday 01-01-2007		Sample ID A2152597	
Age at sample date 18.8		Sample Date 21-10-2025	
Gestational age 12 + 6			
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 12 + 2
PAPP-A	1.98 mIU/mL	0.35	Method CRL Robinson
fb-hCG	39.74 ng/mL	0.97	Scan date 17-10-2025
Risks at sampling date			Crown rump length in mm 59
Age risk 1:1114			Nuchal translucency MoM 0.58
Biochemical T21 risk 1:537			Nasal bone present
Combined trisomy 21 risk 1:3500			Sonographer N A
Trisomy 13/18 + NT <1:10000			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 3500 women with the same data, there is one woman with a trisomy 21 pregnancy and 3499 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