

Prisca 5.1.0.17
Date of report: 06-01-2026

Patient data			
Name Mrs. S.SWARNA W/O. YAADAGIRI		Patient ID 0312601050045	
Birthday 12-08-2000		Sample ID A0792825	
Age at sample date 25.4		Sample Date 05-01-2026	
Gestational age 11 + 6			
Correction factors			
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 58	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 11 + 6
PAPP-A	1.56 mIU/mL	0.50	Method CRL Robinson
fb-hCG	45.95 ng/mL	0.94	Scan date 05-01-2026
Risks at sampling date			Crown rump length in mm 54
Age risk 1:921			Nuchal translucency MoM 1.53
Biochemical T21 risk 1:1274			Nasal bone present
Combined trisomy 21 risk 1:836			Sonographer NA
Trisomy 13/18 + NT 1:8200			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 836 women with the same data, there is one woman with a trisomy 21 pregnancy and 835 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 1:8200, which represents a low risk.			

Sign of Physician