

Prisca 5.1.0.17
Date of report: 29-11-2025

Patient data			
Name Mrs. RAJKUMARI YADAV		Patient ID 0382511280125	
Birthday 26-07-1998		Sample ID B3963516	
Age at sample date 27.3		Sample Date 28-11-2025	
Gestational age 12 + 0			
Correction factors			
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 51	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 11 + 6
PAPP-A	2.09 mIU/mL	0.54	Method CRL Robinson
fb-hCG	42.074 ng/mL	0.84	Scan date 27-11-2025
Risks at sampling date			Crown rump length in mm 54
Age risk 1:822			Nuchal translucency MoM 0.70
Biochemical T21 risk 1:1750			Nasal bone present
Combined trisomy 21 risk 1:9959			Sonographer NA
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT Sonographer
Risk			Trisomy 21
The graph illustrates the risk of Trisomy 21. The y-axis shows risk levels from 1:10 to 1:10000. The x-axis shows gestational age from 13 to 49 weeks. A red region at the top indicates risk above the cut-off. A yellow region in the middle indicates risk below the cut-off but above the age risk. A green region at the bottom indicates risk below the cut-off. A black vertical bar at age 27 shows the patient's risk level, which is below the cut-off.			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 9959 women with the same data, there is one woman with a trisomy 21 pregnancy and 9958 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