

Prisca 5.1.0.17
Date of report: 14-05-2025

N A

Patient data			
Name Mrs. ANJALIN BHUSHAN KANDULNA		Patient ID 0662505130258	
Birthday 19-05-1989		Sample ID B2669914	
Age at sample date 36.0		Sample Date 13-05-2025	
Gestational age 11 + 2			
Correction factors			
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 53	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 11 + 2
PAPP-A	1.54 mIU/mL	0.59	Method CRL Robinson
fb-hCG	52.37 ng/mL	0.92	Scan date 13-05-2025
Risks at sampling date			Crown rump length in mm 47.8
Age risk 1:207			Nuchal translucency MoM 0.61
Biochemical T21 risk 1:443			Nasal bone present
Combined trisomy 21 risk 1:2505			Sonographer N A
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT MD
Risk			Trisomy 21
The graph illustrates the calculated risk for Trisomy 21 (with nuchal translucency) compared to the cut-off and age risk. The patient's risk is indicated by a black bar at age 37, which falls within the green region, signifying a low risk.			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 2505 women with the same data, there is one woman with a trisomy 21 pregnancy and 2504 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