

Prisca 5.1.0.17
Date of report: 20-10-2025

N A

Patient data			
Name Mrs. SANGAMITRA PATTANAIK		Patient ID 0652510160065	
Birthday 07-07-1987		Sample ID B3871859	
Age at sample date 38.3		Sample Date 16-10-2025	
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 + 6
PAPP-A	2.74 mIU/mL	0.70	Method CRL Robinson
fb-hCG	46.96 ng/mL	0.94	Scan date 15-10-2025
Risks at sampling date			Crown rump length in mm 53.8
Age risk 1:122			Nuchal translucency MoM 0.35
Biochemical T21 risk 1:386			Nasal bone present
Combined trisomy 21 risk 1:2090			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 2090 women with the same data, there is one woman with a trisomy 21 pregnancy and 2089 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
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