

Prisca 5.1.0.17
Date of report: 18-03-2025

N A

Patient data			
Name Mrs. PRATIKSHA SAGAR KITEY		Patient ID 0372503170059	
Birthday 07-07-2000		Sample ID A1019733	
Age at sample date 24.7		Sample Date 17-03-2025	
Gestational age 11 + 6			
Correction factors			
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 67	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 11 + 5
PAPP-A	0.99 mIU/mL	0.38	Method CRL Robinson
fb-hCG	58.94 ng/mL	1.26	Scan date 16-03-2025
Risks at sampling date			Crown rump length in mm 52
Age risk 1:951			Nuchal translucency MoM 0.57
Biochemical T21 risk 1:310			Nasal bone unknown
Combined trisomy 21 risk 1:2059			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 2059 women with the same data, there is one woman with a trisomy 21 pregnancy and 2058 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

below cut off
 Below Cut Off, but above Age Risk
 above cut off