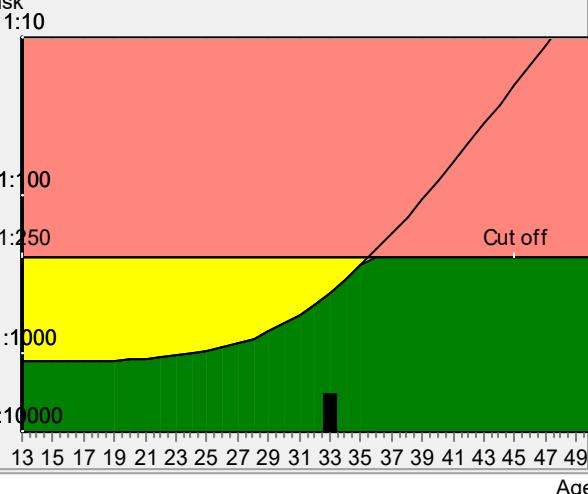


Prisca 5.1.0.17
Date of report: 13/10/25

N A

Patient data								
Name		Mrs. SMITA		Patient ID	0872510110005			
Birthday		22/10/92		Sample ID	A1961847			
Age at sample date		33.0		Sample Date	11/10/25			
Gestational age		13 + 1						
Correction factors								
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	58	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	3.05 mIU/mL	0.58	Gestational age 12 + 6					
fb-hCG	34.41 ng/mL	0.95	Method CRL Robinson					
Risks at sampling date								
Age risk		1:418	Scan date 09/10/25					
Biochemical T21 risk		1:821	Crown rump length in mm 67					
Combined trisomy 21 risk		1:4668	Nuchal translucency MoM 0.56					
Trisomy 13/18 + NT		<1:10000	Nasal bone present					
Risk								
1:10								
								
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 4668 women with the same data, there is one woman with a trisomy 21 pregnancy and 4667 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

 below cut off

 Below Cut Off, but above Age Risk

 above cut off