

Prisca 5.1.0.17

Date of report: 26/09/25

NA

Patient data					
Name	Mrs. SARITA PAWAR	Patient ID	0662509240022		
Birthday	11/06/94	Sample ID	B3401533		
Age at sample date	31.3	Sample Date	24/09/25		
Gestational age	12 + 5				
Correction factors					
Fetuses	1	IVF	no		
Weight	46	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data					
Parameter	Value	Corr. MoM	Ultrasound data		
PAPP-A	4 mIU/mL	0.69	Gestational age 12 + 5		
fb-hCG	39.76 ng/mL	0.90	Method CRL Robinson		
Risks at sampling date					
Age risk	1:545		Scan date 24/09/25		
Biochemical T21 risk	1:1803		Crown rump length in mm 64.8		
Combined trisomy 21 risk	1:8950		Nuchal translucency MoM 0.90		
Trisomy 13/18 + NT	<1:10000		Nasal bone present		
Risk 1:10		Sonographer NA			
1:100		Qualifications in measuring NT NA			
1:250					
1:1000					
1:10000					
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49					
Age					
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 8950 women with the same data, there is one woman with a trisomy 21 pregnancy and 8949 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