

Prisca 5.1.0.17

Date of report: 11/01/25

NA

Patient data			
Name	Mrs. NAMRATA VISHVAKARMA	Patient ID	0622501100050
Birthday	13/12/86	Sample ID	A1630836
Age at sample date	38.1	Sample Date	10/01/25
Gestational age	13 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	71	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	10.19 mIU/mL	2.13	Gestational age 13 + 3
fb-hCG	35.51 ng/mL	1.18	Method CRL Robinson
Risks at sampling date			
Age risk	1:136		Scan date 09/01/25
Biochemical T21 risk	1:2131		Crown rump length in mm 75.9
Combined trisomy 21 risk	1:9161		Nuchal translucency MoM 0.54
Trisomy 13/18 + NT	<1:10000		Nasal bone present
			Sonographer NA
			Qualifications in measuring NT MD
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 9161 women with the same data, there is one woman with a trisomy 21 pregnancy and 9160 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