

Prisca 5.1.0.17

Date of report: 01/12/25

Patient data			
Name	Mrs. SUJATA KUMARI	Patient ID	0522511290055
Birthday	26/03/01	Sample ID	b4164601
Age at sample date	24.7	Sample Date	29/11/25
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	42	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	1.23 mIU/mL	0.17	Gestational age 13 + 0
fb-hCG	37.13 ng/mL	0.88	Method CRL Robinson
Risks at sampling date			
Age risk	1:990		Scan date 29/11/25
Biochemical T21 risk	1:97		Crown rump length in mm 70.6
Combined trisomy 21 risk	1:494		Nuchal translucency MoM 1.01
Trisomy 13/18 + NT	1:525		Nasal bone present
			Sonographer NA
			Qualifications in measuring NT Sonographer
Trisomy 21			
The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. <p>After the result of the Trisomy 21 test (with NT) it is expected that among 494 women with the same data, there is one woman with a trisomy 21 pregnancy and 493 women with not affected pregnancies.</p> <p>The PAPP-A level is low.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.</p> <p>Please note that risk calculations are statistical approaches and have no diagnostic value!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>			
Trisomy 13/18 + NT			
The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:525, which represents a low risk.			

Sign of Physician

below cut off

Below Cut Off, but above Age Risk

above cut off