

Prisca 5.1.0.17
Date of report: 01/12/25

Patient data			
Name	Mrs. SUJATA KUMARI		Patient ID
Birthday	26/03/01	Sample ID	
Age at sample date	24.7	Sample Date	
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	42	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.23 mIU/mL	0.17	
fb-hCG	37.13 ng/mL	0.88	
Risks at sampling date			
Age risk	1:990	Gestational age	13 + 0
Biochemical T21 risk	1:97	Method	CRL Robinson
Combined trisomy 21 risk	1:494	Scan date	29/11/25
Trisomy 13/18 + NT	1:525	Crown rump length in mm	70.6
		Nuchal translucency MoM	1.01
		Nasal bone	present
		Sonographer	NA
		Qualifications in measuring NT	Sonographer
Trisomy 21			
<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <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. 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 hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>			
Trisomy 13/18 + NT			
<p>The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:525, which represents a low risk.</p>			

Sign of Physician