

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
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 12 + 5
PAPP-A	4 mIU/mL	0.69	Method CRL Robinson
fb-hCG	39.76 ng/mL	0.90	Scan date 24/09/25
Risks at sampling date			Crown rump length in mm 64.8
Age risk	1:545		Nuchal translucency MoM 0.90
Biochemical T21 risk	1:1803		Nasal bone present
Combined trisomy 21 risk	1:8950		Sonographer NA
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT NA
Risk			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 8950 women with the same data, there is one woman with a trisomy 21 pregnancy and 8949 women with not affected pregnancies.</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 < 1:10000, which represents a low risk.</p>			

Sign of Physician

below cut off
 Below Cut Off, but above Age Risk
 above cut off