

Prisca 5.1.0.17
Date of report: 28/10/25

NA

Patient data			
Name	Mrs. ARUSHI SINGH	Patient ID	0852510250136
Birthday	22/07/01	Sample ID	B3520121
Age at sample date	24.3	Sample Date	25/10/25
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	52	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.91 mIU/mL	0.45	
fb-hCG	43.35 ng/mL	0.93	
Risks at sampling date			
Age risk		1:982	
Biochemical T21 risk		1:1008	
Combined trisomy 21 risk		1:6121	
Trisomy 13/18 + NT		<1:10000	
			Gestational age 12 + 2
			Method CRL Robinson
			Scan date 25/10/25
			Crown rump length in mm 60
			Nuchal translucency MoM 0.64
			Nasal bone present
			Sonographer NA
			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 6121 women with the same data, there is one woman with a trisomy 21 pregnancy and 6120 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