

Prisca 5.1.0.17
Date of report: 11-09-2025

N A

Patient data			
Name	Mrs. POOJA SHARMA		Patient ID
Birthday	06-09-1993	Sample ID	0622509100064 A1827232
Age at sample date	32.0	Sample Date	10-09-2025
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	86	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.88 mIU/mL	1.14	12 + 4
fb-hCG	33.1 ng/mL	1.07	Method
			CRL Robinson
			Scan date
			05-09-2025
			Crown rump length in mm
			63.7
			Nuchal translucency MoM
			0.86
			Nasal bone
			present
			Sonographer
			N A
			Qualifications in measuring NT
			MD
Risks at sampling date			
Age risk	1:497		
Biochemical T21 risk	1:3520		
Combined trisomy 21 risk	<1:10000		
Trisomy 13/18 + NT	<1:10000		
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</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