

Prisca 5.1.0.17
Date of report: 20-05-2025

N A

Patient data				
Name	Mrs. HARSHALI SHELAR		Patient ID	0662505180101
Birthday	10-05-1996		Sample ID	B2665145
Age at sample date	29.0		Sample Date	18-05-2025
Gestational age	12 + 1			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	50	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 0
PAPP-A	3.52 mIU/mL	0.84	Method	CRL Robinson
fb-hCG	44.52 ng/mL	0.92	Scan date	17-05-2025
Risks at sampling date			Crown rump length in mm	
Age risk	1:712		56	
Biochemical T21 risk	1:3695		Nuchal translucency MoM	
Combined trisomy 21 risk	<1:10000		0.95	
Trisomy 13/18 + NT	<1:10000		Nasal bone	
			present	
			Sonographer	
			N A	
			Qualifications in measuring NT	
			MD	
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