

Prisca 5.1.0.17
Date of report: 27/12/25

Patient data			
Name	Mrs. RANJANA MOHURLE		Patient ID
Birthday	15/04/99	Sample ID	
Age at sample date	26.7	Sample Date	
Gestational age	13 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	40	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.48 mIU/mL	0.48	
fb-hCG	32.96 ng/mL	0.89	
Risks at sampling date			
Age risk	1:908		
Biochemical T21 risk	1:1236		
Combined trisomy 21 risk	1:7329		
Trisomy 13/18 + NT	<1:10000		
		Gestational age	13 + 2
		Method	CRL Robinson
		Scan date	24/12/25
		Crown rump length in mm	73
		Nuchal translucency MoM	0.50
		Nasal bone	present
		Sonographer	NA
		Qualifications in measuring NT	Sonographer
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 7329 women with the same data, there is one woman with a trisomy 21 pregnancy and 7328 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