

Prisca 5.1.0.17
Date of report: 06/10/25

NA

Patient data			
Name	Mrs. NAYLA PARWEEN	Patient ID	0482509280053
Birthday	01/01/94	Sample ID	B3485111
Age at sample date	31.7	Sample Date	28/09/25
Gestational age	11 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	70	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	0.22 mIU/mL	0.09	
fb-hCG	48.08 ng/mL	1.01	
Risks at sampling date			
Age risk	1:489	Gestational age	11 + 5
Biochemical T21 risk	>1:50	Method	CRL Robinson
Combined trisomy 21 risk	>1:50	Scan date	28/09/25
Trisomy 13/18 + NT	>1:50	Crown rump length in mm	51.6
		Nuchal translucency MoM	1.66
		Nasal bone	present
		Sonographer	NA
		Qualifications in measuring NT	NA
Risk		Trisomy 21	
		The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk. After the result of the Trisomy 21 Test (with nuchal translucency), it is expected that among less than 50 pregnancies with the same data, there is one trisomy 21 pregnancy. The PAPP-A level is low. 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! The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)). The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
Trisomy 13/18 + NT			
The calculated risk for Trisomy 13/18 test (with nuchal translucency) is >1:50, which indicates an increased risk.			

Sign of Physician

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