

Prisca 5.1.0.17
Date of report: 30-10-2025

NA

Patient data			
Name	Mrs. MANTSHA	Patient ID	0772510280119
Birthday	01-01-1998	Sample ID	B3669105
Age at sample date	27.8	Sample Date	28-10-2025
Gestational age	12 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	43.6	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.16 mIU/mL	0.69	11 + 4
fb-hCG	46.15 ng/mL	0.87	Method
			CRL Robinson
			Scan date
			25-10-2025
Risks at sampling date			Trisomy 21
Age risk	1:792		Crown rump length in mm
Biochemical T21 risk	1:2843		51.3
Combined trisomy 21 risk	>1:50		Nuchal translucency MoM
Trisomy 13/18 + NT	>1:50		3.70
			Nasal bone
			unknown
			Sonographer
			NA
			Qualifications in measuring NT
			NA
Risk Age 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 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