

Prisca 5.1.0.17
Date of report: 27/12/25

Patient data			
Name	Mrs. MANISHA KUMARI	Patient ID	0472512260073
Birthday	08/02/00	Sample ID	B3854836
Age at sample date	25.9	Sample Date	26/12/25
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	44	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.52 mIU/mL	0.44	
fb-hCG	32.96 ng/mL	0.89	
Risks at sampling date			
Age risk	1:949	Gestational age	13 + 2
Biochemical T21 risk	1:1029	Method	CRL Robinson
Combined trisomy 21 risk	1:6238	Scan date	25/12/25
Trisomy 13/18 + NT	<1:10000	Crown rump length in mm	73.1
		Nuchal translucency MoM	0.61
		Nasal bone	present
		Sonographer	NA
		Qualifications in measuring NT	Sonographer
Risk		Trisomy 21	
		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 6238 women with the same data, there is one woman with a trisomy 21 pregnancy and 6237 women with not affected pregnancies. 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 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician

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