

Prisca 5.1.0.17
Date of report: 13/12/25

Patient data			
Name	Mrs. SUMAN DAS	Patient ID	0692512110217
Birthday	01/06/95	Sample ID	B4105099
Age at sample date	30.5	Sample Date	11/12/25
Gestational age	11 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	64	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.52 mIU/mL	0.78	
fb-hCG	50.01 ng/mL	0.91	
Risks at sampling date			
Age risk	1:571	Gestational age	11 + 1
Biochemical T21 risk	1:2490	Method	CRL Robinson
Combined trisomy 21 risk	1:9506	Scan date	11/12/25
Trisomy 13/18 + NT	<1:10000	Crown rump length in mm	46.1
		Nuchal translucency MoM	1.03
		Nasal bone	present
		Sonographer	NA
		Qualifications in measuring NT	Sonographer
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 9506 women with the same data, there is one woman with a trisomy 21 pregnancy and 9505 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 held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!			
Risk 1:10 1:100 1:250 1:1000 1:10000 Age			
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