

Prisca 5.1.0.17
Date of report: 22-12-2024

NA

Patient data			
Name	Mrs. SANGAM KUMARI		Patient ID
Birthday	21-01-1997	Sample ID	0622412200019
Age at sample date	27.9	Sample Date	25165033
Gestational age	13 + 1		20-12-2024
Correction factors			
Fetuses	1	IVF	no
Weight	68	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	4.1 mIU/mL	0.95	12 + 6
fb-hCG	24.3 ng/mL	0.71	Method
			CRL Robinson
			Scan date
			18-12-2024
			Crown rump length in mm
			68
			Nuchal translucency MoM
			0.94
			Nasal bone
			absent
			Sonographer
			NA
			Qualifications in measuring NT
			MD
Risks at sampling date		Trisomy 21	
Age risk	1:818	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 842 women with the same data, there is one woman with a trisomy 21 pregnancy and 841 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!	
Biochemical T21 risk	1:9673		
Combined trisomy 21 risk	1:842		
Trisomy 13/18 + NT	<1:10000		
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