

Prisca 5.1.0.17
Date of report: 11-12-2024

NA

Patient data			
Name	Mrs. SARIKA SUBHASH SAHARE		Patient ID
Birthday	28-10-1989	Sample ID	A1017855
Age at sample date	35.1	Sample Date	10-12-2024
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	52.5	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.61 mIU/mL	0.61	Method
fb-hCG	35.25 ng/mL	0.94	Scan date
			Crown rump length in mm
			Nuchal translucency MoM
			Nasal bone
			Sonographer
			Qualifications in measuring NT
Risks at sampling date		Trisomy 21	
Age risk	1:271	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 3400 women with the same data, there is one woman with a trisomy 21 pregnancy and 3399 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	1:622		
Combined trisomy 21 risk	1:3400		
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