

Prisca 5.1.0.17
Date of report: 18-09-2024

N A

Patient data			
Name	Mrs. SARLA SATNAMI		Patient ID
Birthday	10-06-1989	Sample ID	a1047936
Age at sample date	35.3	Sample Date	16-09-2024
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	55	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	5.39 mIU/mL	1.02	13 + 0
fb-hCG	51.1 ng/mL	1.33	Method
			CRL Robinson
			Scan date
			16-09-2024
Risks at sampling date			Crown rump length in mm
Age risk	1:261		69
Biochemical T21 risk	1:872		Nuchal translucency MoM
Combined trisomy 21 risk	1:1529		1.27
Trisomy 13/18 + NT	<1:10000		Nasal bone
			unknown
			Sonographer
			N A
			Qualifications in measuring NT
			MD
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 1529 women with the same data, there is one woman with a trisomy 21 pregnancy and 1528 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