

Prisca 5.1.0.17
Date of report: 14-01-2025

NA

Patient data			
Name	Mrs. SUJITHA	Patient ID	0042501130003
Birthday	17-10-2005	Sample ID	23125759
Age at sample date	19.2	Sample Date	13-01-2025
Gestational age	12 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	37	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.84 mIU/mL	0.48	12 + 1
fb-hCG	161.26 ng/mL	2.94	Method
			CRL Robinson
			Scan date
			13-01-2025
Risks at sampling date		Trisomy 21	
Age risk	1:1083	Crown rump length in mm	57.1
Biochemical T21 risk	1:85	Nuchal translucency MoM	0.67
Combined trisomy 21 risk	1:596	Nasal bone	present
Trisomy 13/18 + NT	<1:10000	Sonographer	NA
		Qualifications in measuring NT	MD
		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 596 women with the same data, there is one woman with a trisomy 21 pregnancy and 595 women with not affected pregnancies. The free beta HCG level is high. 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