

Patient data				
Name	Mrs. N.BHAGYA LAXMI		Patient ID	0352208290049
Birthday	20-12-1991		Sample ID	m1429446
Age at sample date	30.7		Sample Date	29-08-2022
Gestational age	12 + 5			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	63	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 3
PAPP-A	3.15 mIU/mL	0.78	Method	CRL Robinson
fb-hCG	45.66 ng/mL	1.16	Scan date	27-08-2022
Risks at sampling date			Crown rump length in mm	
Age risk	1:594		61	
Biochemical T21 risk	1:1517		Nuchal translucency MoM	
Combined trisomy 21 risk	1:192		1.90	
Trisomy 13/18 + NT	1:6066		Nasal bone	
			unknown	
			Sonographer	
			NA	
			Qualifications in measuring NT	
			NAGA SAI TEJA	
Risk			Trisomy 21	
			The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk. After the result of the Trisomy 21 test (with NT) it is expected that among 192 women with the same data, there is one woman with a trisomy 21 pregnancy and 191 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 1:6066, which represents a low risk.				

Sign of Physician