

Patient data					
Name	Ms. T SHANTHA KUMARI		Patient ID	0012208270602	
Birthday	25-06-1997		Sample ID	23590427	
Age at sample date	25.2		Sample Date	27-08-2022	
Gestational age	11 + 5				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	45	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	11 + 6	
PAPP-A	1.79 mIU/mL	0.46	Method	CRL Robinson	
fb-hCG	49.65 ng/mL	0.89	Scan date	28-08-2022	
Risks at sampling date			Crown rump length in mm	55	
Age risk	1:926		Nuchal translucency MoM	0.82	
Biochemical T21 risk	1:1110		Nasal bone	unknown	
Combined trisomy 21 risk	1:6568		Sonographer	NA	
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	NAGA SAI TEJA	
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 6568 women with the same data, there is one woman with a trisomy 21 pregnancy and 6567 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