

Patient data			
Name	Mrs. ASHWINI	Patient ID	0352207300083
Birthday	19-05-1989	Sample ID	23624588
Age at sample date	33.2	Sample Date	30-07-2022
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	69	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.62 mIU/mL	0.50	
fb-hCG	45.22 ng/mL	1.10	
Risks at sampling date			
Age risk		1:391	
Biochemical T21 risk		1:373	
Combined trisomy 21 risk		1:2251	
Trisomy 13/18 + NT		<1:10000	
			Gestational age 11 + 5
			Method CRL Robinson
			Scan date 25-07-2022
			Crown rump length in mm 52.9
			Nuchal translucency MoM 0.71
			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 below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 2251 women with the same data, there is one woman with a trisomy 21 pregnancy and 2250 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 held 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