

Patient data			
Name	Mrs. M BHARGAVI	Patient ID	0012303020620
Birthday	26-01-2004	Sample ID	23913138
Age at sample date	19.1	Sample Date	02-03-2023
Gestational age	12 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	58	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.56 mIU/mL	1.08	12 + 0
fb-hCG	51.22 ng/mL	1.08	Method
			CRL Robinson
			Scan date
			02-03-2023
Risks at sampling date			Crown rump length in mm
Age risk	1:1079		56
Biochemical T21 risk	1:6674		Nuchal translucency MoM
Combined trisomy 21 risk	<1:10000		0.68
Trisomy 13/18 + NT	<1:10000		Nasal bone
			unknown
			Sonographer
			NA
			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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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