

Patient data				
Name	Mrs. G.MAHANVITHA		Patient ID	0352206100089
Birthday	06/05/99		Sample ID	22664870
Age at sample date	23.1		Sample Date	10/06/22
Gestational age	11 + 6			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	64	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	11 + 2
PAPP-A	1.23 mIU/mL	0.45	Method	CRL Robinson
fb-hCG	49.66 ng/mL	1.04	Scan date	06/06/22
Risks at sampling date			Crown rump length in mm	47.3
Age risk	1:1003		Nuchal translucency MoM	0.70
Biochemical T21 risk	1:785		Nasal bone	unknown
Combined trisomy 21 risk	1:4856		Sonographer	DR NA
Trisomy 13/18 + NT	<1:10000		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 4856 women with the same data, there is one woman with a trisomy 21 pregnancy and 4855 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