

Patient data				
Name	Mrs. K LAVANYA W/O SRINIVAS		Patient ID	0012206160344
Birthday	10/12/96		Sample ID	23534255
Age at sample date	25.5		Sample Date	16/06/22
Gestational age	13 + 1			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	50	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	13 + 0
PAPP-A	1.55 mIU/mL	0.25	Method	CRL Robinson
fb-hCG	35.69 ng/mL	0.94	Scan date	15/06/22
Risks at sampling date			Crown rump length in mm	70
Age risk	1:958		Nuchal translucency MoM	1.76
Biochemical T21 risk	1:169		Nasal bone	unknown
Combined trisomy 21 risk	>1:50		Sonographer	DR NA
Trisomy 13/18 + NT	1:126		Qualifications in measuring NT	MD
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 nuchal translucency), it is expected that among less than 50 pregnancies with the same data, there is one trisomy 21 pregnancy. The PAPP-A level is low. 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 1:126, which represents a low risk.				

 Sign of Physician