

Patient data			
Name		Mrs. B RAMYA	
Patient ID		0352211020021	
Birthday		22-10-2002	
Sample ID		23745479	
Age at sample date		20.0	
Sample Date		31-10-2022	
Gestational age		13 + 6	
Correction factors			
Fetuses	1	IVF	no
Weight	50	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	12.65 mIU/mL	1.59	
fb-hCG	34.21 ng/mL	1.10	
Gestational age		13 + 5	
Method		CRL Robinson	
Scan date		30-10-2022	
Risks at sampling date		Crown rump length in mm	
Age risk		1:1135	
Biochemical T21 risk		<1:10000	
Nuchal translucency MoM		0.62	
Combined trisomy 21 risk		<1:10000	
Nasal bone		unknown	
Trisomy 13/18 + NT		<1:10000	
Sonographer		NA	
Qualifications in measuring NT		NA	
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

below cut off
 Below Cut Off, but above Age Risk
 above cut off