

Patient data			
Name	Mrs. G.JHANSI	Patient ID	0352407160016
Birthday	02-01-2005	Sample ID	23499464
Age at sample date	19.5	Sample Date	15-07-2024
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	59	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.87 mIU/mL	0.80	
fb-hCG	32.14 ng/mL	0.86	
Risks at sampling date			Gestational age
Age risk	1:1112		13 + 0
Biochemical T21 risk	1:5804		Method
Combined trisomy 21 risk	<1:10000		CRL Robinson
Trisomy 13/18 + NT	<1:10000		Scan date
			15-07-2024
			Crown rump length in mm
			70
			Nuchal translucency MoM
			0.74
			Nasal bone
			present
			Sonographer
			N A
			Qualifications in measuring NT
			MD
Risk		Trisomy 21	
1:10		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!	
1:100			
1:250			
1:1000			
1:10000			
1:10000			
Age			
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