

Patient data			
Name		Mrs. G JYOTHI	
Patient ID		0012508050096	
Birthday		15/08/99	
Sample ID		A1907730	
Age at sample date		26.0	
Sample Date		05/08/25	
Gestational age		13 + 2	
Correction factors			
Fetuses	1	IVF	no
Weight	72	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.01 mIU/mL	0.24	13 + 1
fb-hCG	36.85 ng/mL	1.13	Method
Risks at sampling date			CRL Robinson
Age risk	1:940		Scan date
Biochemical T21 risk	1:93		04/08/25
Combined trisomy 21 risk	1:696		Crown rump length in mm
Trisomy 13/18 + NT	1:6504		72
Risk			Nuchal translucency MoM
1:10			0.67
1:100			Nasal bone
1:250			present
1:1000			Sonographer
1:10000			N A
Age			Qualifications in measuring NT
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			MD
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 696 women with the same data, there is one woman with a trisomy 21 pregnancy and 695 women with not affected pregnancies. 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 1:6504, which represents a low risk.			

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