

Patient data			
Name		Mrs. VASANTHA	
Patient ID		0012409250395	
Birthday		30-03-2001	
Sample ID		a1301813	
Age at sample date		23.5	
Sample Date		25-09-2024	
Gestational age		13 + 3	
Correction factors			
Fetuses	1	IVF	no
Weight	60	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.69 mIU/mL	1.02	
fb-hCG	25.33 ng/mL	0.77	
Risks at sampling date			
Age risk	1:1047		
Biochemical T21 risk	<1:10000		
Combined trisomy 21 risk	<1:10000		
Trisomy 13/18 + NT	<1:10000		
			Gestational age 12 + 6 Method CRL Robinson Scan date 21-09-2024 Crown rump length in mm 67 Nuchal translucency MoM 0.82 Nasal bone present Sonographer N A Qualifications in measuring NT MD
Risk 1:10 1:100 1:250 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age			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.			

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