

Patient data			
Name	Mrs. SRIVIDYA	Patient ID	0352504080051
Birthday	10/06/98	Sample ID	B2410972
Age at sample date	26.8	Sample Date	08/04/25
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	42	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.16 mIU/mL	0.44	Gestational age
fb-hCG	35.74 ng/mL	0.84	Method
Risks at sampling date			CRL Robinson
Age risk	1:884		Scan date
Biochemical T21 risk	1:1062		Crown rump length in mm
Combined trisomy 21 risk	1:6150		Nuchal translucency MoM
Trisomy 13/18 + NT	<1:10000		Nasal bone
			present
			Sonographer
			N A
			Qualifications in measuring NT
			MD
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 6150 women with the same data, there is one woman with a trisomy 21 pregnancy and 6149 women with not affected pregnancies. 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