

N A

Patient data					
Name	Mrs. SRILEKHA POLURU		Patient ID	0312408110050	
Birthday	26-08-1992		Sample ID	a0828819	
Age at sample date	32.0		Sample Date	11-08-2024	
Gestational age	13 + 2				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	64	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	13 + 2	
PAPP-A	4.75 mIU/mL	0.97	Method	CRL Robinson	
fb-hCG	53.69 ng/mL	1.59	Scan date	11-08-2024	
Risks at sampling date			Crown rump length in mm	75	
Age risk	1:501		Nuchal translucency MoM	0.87	
Biochemical T21 risk	1:983		Nasal bone	present	
Combined trisomy 21 risk	1:5001		Sonographer	N A	
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	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 5001 women with the same data, there is one woman with a trisomy 21 pregnancy and 5000 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.					

Sign of Physician

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