

NA

Patient data					
Name	Mrs. PARVATHI		Patient ID	0352411230020	
Birthday	10-02-1986		Sample ID	A1251532	
Age at sample date	38.8		Sample Date	23-11-2024	
Gestational age	13 + 5				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21	unknown
Weight	56	diabetes	no	pregnancies	
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	13 + 2	
PAPP-A	8.69 mIU/mL	1.07	Method	CRL Robinson	
fb-hCG	36.54 ng/ml	1.18	Scan date	20-11-2024	
Risks at sampling date			Crown rump length in mm	75	
Age risk	1:114		Nuchal translucency MoM	0.81	
Biochemical T21 risk	1:565		Nasal bone	present	
Combined trisomy 21 risk	1:2825		Sonographer	NA	
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 2825 women with the same data, there is one woman with a trisomy 21 pregnancy and 2824 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