

N A

Patient data			
Name	Mrs. JHIMLI ROUTH		Patient ID
Birthday	13/01/90	Sample ID	A0982183
Age at sample date	34.6	Sample Date	12/08/24
Gestational age	11 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	49	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.6 mIU/mL	1.22	
fb-hCG	50.31 ng/mL	0.97	
Risks at sampling date			
Age risk	1:290		Gestational age
Biochemical T21 risk	1:2955		Method
Combined trisomy 21 risk	1:8453		Scan date
Trisomy 13/18 + NT	<1:10000		Crown rump length in mm
			50
			Nuchal translucency MoM
			1.11
			Nasal bone
			unknown
			Sonographer
			N A
			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 8453 women with the same data, there is one woman with a trisomy 21 pregnancy and 8452 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