

Patient data			
Name	Mrs. JAGRITI PRIYA	Patient ID	0472310280079
Birthday	02/10/96	Sample ID	24909868
Age at sample date	27.1	Sample Date	28/10/23
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	58	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.08 mIU/mL	0.78	
fb-hCG	40.41 ng/mL	1.12	
Risks at sampling date			
Age risk		1:873	
Biochemical T21 risk		1:2434	
Combined trisomy 21 risk		1:9408	
Trisomy 13/18 + NT		<1:10000	
			Gestational age 13 + 0
			Method CRL Robinson
			Scan date 27/10/23
			Crown rump length in mm 70
			Nuchal translucency MoM 1.02
			Nasal bone present
			Sonographer MANISHA CHOUDHARY
			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 9408 women with the same data, there is one woman with a trisomy 21 pregnancy and 9407 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