

TRIVENI MS(O,B,G)

Patient data					
Name	Mrs. PRATHYUSHA W/O SUNIL		Patient ID	0312305220053	
Birthday	26-08-1996		Sample ID	23981933	
Age at sample date	26.7		Sample Date	22-05-2023	
Gestational age	13 + 3				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	58	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	11 + 5	
PAPP-A	2.17 mIU/mL	0.37	Method	CRL Robinson	
fb-hCG	31.8 ng/mL	0.95	Scan date	10-05-2023	
Risks at sampling date			Crown rump length in mm	52	
Age risk	1:902		Nuchal translucency MoM	1.01	
Biochemical T21 risk	1:530		Nasal bone	present	
Combined trisomy 21 risk	1:2435		Sonographer	TRIVENI MS(O,B,G)	
Trisomy 13/18 + NT	<1:10000		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 2435 women with the same data, there is one woman with a trisomy 21 pregnancy and 2434 women with not affected pregnancies. The PAPP-A level is low. 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