

Prisca

5.1.0.17

Date of report:

27-11-2024

NA

Patient data			
Name	Mrs. SARITA GARG FETUS-B		Patient ID
Birthday	24-01-1992	Sample ID	A1398757 FETUS-B
Age at sample date	32.8	Sample Date	26-11-2024
Gestational age	12 + 5		
Correction factors			
Fetuses	2	IVF	no
Weight	58.1	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	12.65 mIU/mL	1.37	Gestational age
fb-hCG	131.67 ng/ml	1.59	Method
			CRL Robinson
			Scan date
			25-11-2024
			Crown rump length in mm
			64
			Nuchal translucency MoM
			0.73
			Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			MD
Risks at sampling date			Trisomy 21
Age risk	1:422		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 8115 women with the same data, there is one woman with a trisomy 21 pregnancy and 8114 women with not affected pregnancies. The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs. 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!
Biochemical T21 risk	1:1652		
Combined trisomy 21 risk	1:8115		
Trisomy 13/18 + NT	<1:10000		
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