

Prisca

5.1.0.17

Date of report:

30/03/21

Patient data			
Name	Mrs. LAXMI Fetus-A		Patient ID
Birthday	10/02/90	Sample ID	0012103280042
Age at sample date	31.1	Sample Date	S1066354
Gestational age	11 + 5		27/03/21
Correction factors			
Fetuses	2	IVF	no
Weight	66	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.01 mIU/mL	0.43	11 + 4
fb-hCG	56.98 ng/mL	0.54	Method
			CRL Robinson
			Scan date
			26/03/21
			Crown rump length in mm
			49.95
			Nuchal translucency MoM
			0.89
			Nasal bone
			unknown
			Sonographer
			DR NA
			Qualifications in measuring NT
			MD
Risks at sampling date		Trisomy 21	
Age risk	1:538	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 8226 women with the same data, there is one woman with a trisomy 21 pregnancy and 8225 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	1:1537		
Combined trisomy 21 risk	1:8226		
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