

Prisca

5.1.0.17

Date of report:

12/05/21

Patient data			
Name	Ms. K DEEPIKA Fetus-B		Patient ID
Birthday	08/06/89	Sample ID	19019050
Age at sample date	31.9	Sample Date	10/05/21
Gestational age	11 + 5		
Correction factors			
Fetuses	2	IVF	no
Weight	59	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.33 mIU/mL	0.63	
fb-hCG	56.15 ng/mL	0.52	
Risks at sampling date			
Age risk		1:475	
Biochemical T21 risk		1:4040	
Combined trisomy 21 risk		1:7532	
Trisomy 13/18 + NT		<1:10000	
			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 7532 women with the same data, there is one woman with a trisomy 21 pregnancy and 7531 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!
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