

Prisca 5.1.0.17
Date of report: 15/09/24

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Patient data			
Name	Mrs. TEJASWINI		Patient ID
Birth day	19/04/98	Sample ID	
Age at sample date	26.4	Sample Date	
Gestational age	11 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	30	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.32 mIU/mL	0.48	
fb-hCG	49.5 ng/mL	0.78	
Risks at sampling date			
Age risk	1:862		
Biochemical T21 risk	1:1579		
Combined trisomy 21 risk	1:9123		
Trisomy 13/18 + NT	<1:10000		
			Gestational age
			Method
			Scan date
			Crown rump length in mm
			Nuchal translucency MoM
			Nasal bone
			Sonographer
			Qualifications in measuring NT
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 9123 women with the same data, there is one woman with a trisomy 21 pregnancy and 9122 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

below cut off
 Below Cut Off, but above Age Risk
 above cut off