

Prisca 5.1.0.17
Date of report: 04-10-2024

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Patient data			
Name	Mrs. PRIYANKA ROHANKAR		Patient ID
Birthday	09-07-1992		Sample ID
Age at sample date	32.2		Sample Date
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	58	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	4.1 mIU/mL	0.83	Method
fb-hCG	56.29 ng/mL	1.50	Scan date
Risks at sampling date			Crown rump length in mm
Age risk	1:474		Nuchal translucency MoM
Biochemical T21 risk	1:758		Nasal bone
Combined trisomy 21 risk	1:2667		Sonographer
Trisomy 13/18 + NT	<1:10000		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 2667 women with the same data, there is one woman with a trisomy 21 pregnancy and 2666 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