

Prisca 5.1.0.17
Date of report: 19-10-2025

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Patient data			
Name	Mrs. PINKY PATIAR	Patient ID	0652510170193
Birthday	22-05-2002	Sample ID	B3872679
Age at sample date	23.4	Sample Date	17-10-2025
Gestational age	11 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	41	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.66 mIU/mL	0.97	
fb-hCG	48.52 ng/mL	0.80	
Risks at sampling date			
Age risk		1:978	
Biochemical T21 risk		1:9493	
Combined trisomy 21 risk		1:8298	
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 8298 women with the same data, there is one woman with a trisomy 21 pregnancy and 8297 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