

Prisca 5.1.0.17
Date of report: 23-01-2025

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Patient data			
Name	Mrs. PAYAL CHORGHE		Patient ID
Birthday	02-08-2002	Sample ID	A2090232
Age at sample date	22.5	Sample Date	22-01-2025
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	56	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.28 mIU/mL	0.75	12 + 3
fb-hCG	84.2 ng/mL	1.99	Method
			CRL Robinson
			Scan date
			21-01-2025
Risks at sampling date			Trisomy 21
Age risk	1:1046		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
Biochemical T21 risk	1:664		After the result of the Trisomy 21 test (with NT) it is expected that among 1565 women with the same data, there is one woman with a trisomy 21 pregnancy and 1564 women with not affected pregnancies.
Combined trisomy 21 risk	1:1565		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!
Trisomy 13/18 + NT	<1:10000		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