

Prisca 5.1.0.17
Date of report: 01-05-2025

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Patient data			
Name	Mrs. AROHI	Patient ID	0772504290091
Birthday	01-01-1995	Sample ID	A1960675
Age at sample date	30.3	Sample Date	29-04-2025
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	66.15	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.75 mIU/mL	0.52	12 + 2
fb-hCG	41.12 ng/mL	0.99	Method
			CRL Robinson
			Scan date
			28-04-2025
Risks at sampling date			Trisomy 21
Age risk		1:618	The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
Biochemical T21 risk		1:803	After the result of the Trisomy 21 test (with NT) it is expected that among 517 women with the same data, there is one woman with a trisomy 21 pregnancy and 516 women with not affected pregnancies.
Combined trisomy 21 risk		1:517	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		1:6705	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!
Age			
Trisomy 13/18 + NT			
The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:6705, which represents a low risk.			

Sign of Physician