

Prisca 5.1.0.17
Date of report: 17-10-2024

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Patient data			
Name	Mrs. SHITAL GORE	Patient ID	0662410140161
Birthday	01-10-1984	Sample ID	A1345241
Age at sample date	40.0	Sample Date	14-10-2024
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	yes
Weight	62	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	5.99 mIU/mL	1.54	12 + 6
fb-hCG	43.69 ng/mL	1.07	Method
			CRL Robinson
			Scan date
			16-10-2024
Risks at sampling date			Trisomy 21
Age risk		1:78	The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
Biochemical T21 risk		1:991	After the result of the Trisomy 21 test (with NT) it is expected that among 2367 women with the same data, there is one woman with a trisomy 21 pregnancy and 2366 women with not affected pregnancies.
Combined trisomy 21 risk		1:2367	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