

Prisca 5.1.0.17
Date of report: 29-09-2024

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Patient data			
Name	Mrs. POOJA DANGI	Patient ID	0442409270036
Birthday	20-07-1993	Sample ID	A1041627
Age at sample date	31.2	Sample Date	27-09-2024
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	54	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.63 mIU/mL	0.61	12 + 3
fb-hCG	46.39 ng/mL	1.05	Method
			CRL Robinson
			Scan date
			27-09-2024
Risks at sampling date			Trisomy 21
Age risk		1:548	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 4644 women with the same data, there is one woman with a trisomy 21 pregnancy and 4643 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!
Biochemical T21 risk		1:973	
Combined trisomy 21 risk		1:4644	
Trisomy 13/18 + NT		<1:10000	
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