Molecular Quest Healthcare Pvt. Ltd Plot No. 28-29,Sector 18(P), Electronic City, Gurgaon-122 015 (Haryana)

Prisca 5.2.0.13

Date of report: 9/23/2023

Dr.

Patient data				
Name	MRS SHABNAM		Patient ID	
Birthday	7/1/1994	Sample ID		11803248
Age at sample date	29.2	Sample Date		9/20/2023
Gestational age	12 + 3			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	unknown
Weight 52	diabetes	no		
Smoker no	Origin	Asian		
Biochemical data	Ultrasound data			
Parameter Value	Corr. MoM Gestation		age	12 + 3
PAPP-A 12 mIU/m	nl 1.94	Method CRL Robinson		
fb-hCG 66.8 ng/ml	1.66	Scan date 9/20/2023		
Risks at sampling date			Crown rump length in mm	
Age risk	1:704		Nuchal translucency MoM	
Biochemical T21 risk	1:4608	Nasal bone		unknown
Combined trisomy 21 risk	ny 21 risk <1:10000		Sonographer	
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT		M.D
1:100 1:250 1:1000 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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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!			



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