

Date of Report 11-10-2023  
PRISCA 5.2.0.13

Patient Data			
Name	MRS.BABITA	Patient ID	012310100216
Birthday	20-01-2000	Sample ID	11805664
Age at Sample date	23.7	Sample Date	10-10-2023
Gestational age	13+1		

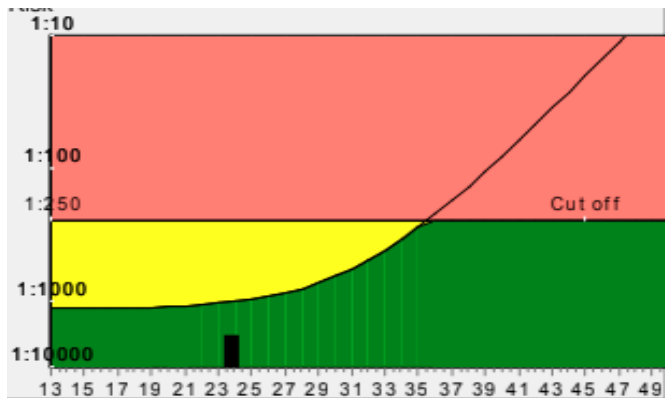
Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	58	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
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Parameter	Value	Corr Mom		
PAPP-A	4.38 mIU/ml	0.62	Gestational age	12+4
fb-hCG	38.4 ng/ml	1.21	Method	CRL (<>Robinson)
			Scan date	06-10-2023

Risks at sampling date				
Age Risk		1:1030	Crown rump length in mm	61.1
Biochemical T21 risk		1:1353	Nuchal translucency MoM	0.89
Combined trisomy 21 risk		1:7191	Nasal bone	PRESENT
Trisomy 13/18 + NT		<1:10000	Sonographer	DR
			Qualifications in measuring NT	

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
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The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 7191 women with the same data, there is one woman with a trisomy 21 pregnancy and 7190 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 the risk calculations are statistical approaches and have no diagnostic value!

The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).

The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

The laboratory cannot be held responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

Risk Above Cut Off
  Risk above Age Risk
  Risk below Age risk