

Date of Report 30-09-2023
PRISCA 5.2.0.13

Patient Data	
Name MRS KARISHMA	Patient ID 012309290161
Birthday 18-08-1996	Sample ID 11781105
Age at Sample date 27.1	Sample Date 03-09-2023
Gestational age 11+5	

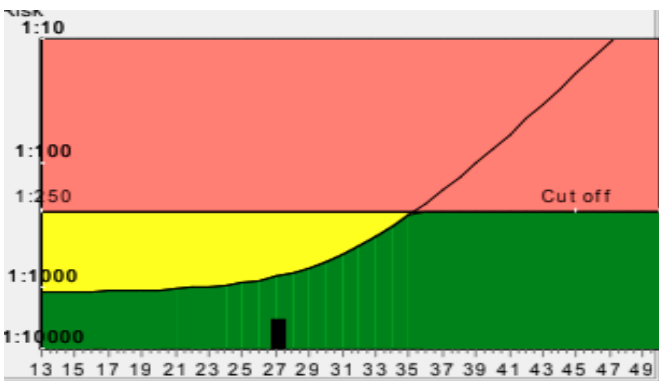
Correction factors			
Fetuses 1	IVF unknown	Previous trisomy 21 unknown	
Weight in kg 48	Diabetes NO	Pregnancies unknown	
Smoker NO	Origin Asian		

Biochemical Data	Ultrasound Data
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Parameter	Value	Corr Mom	
PAPP-A	4.14 mIU/ml	0.82	Method CRL (<>Robinson)
fb-hCG	60.7 ng/ml	1.23	Scan date 29-09-2023

Risks at sampling date	Ultrasound Data
Age Risk 1:827	Crown rump length in mm 49.6
Biochemical T21 risk 1:2073	Nuchal translucency MoM 1.04
Combined trisomy 21 risk 1:7568	Nasal bone PRESENT
Trisomy 13/18 + NT <1:10000	Sonographer DR VIPUL GOYAL
	Qualifications in measuring NT MD

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 7568 women with the same data, there is one woman with a trisomy 21 pregnancy and 7567 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 hold 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