

Date of Report 09-01-2024
PRISCA 5.2.0.13

Patient Data	
Name MRS KIRAN KANWR	Patient ID 012401080115
Birthday 19-01-2003	Sample ID 11827646
Age at Sample date 21.0	Sample Date 08-01-2024
Gestational age 13+6	

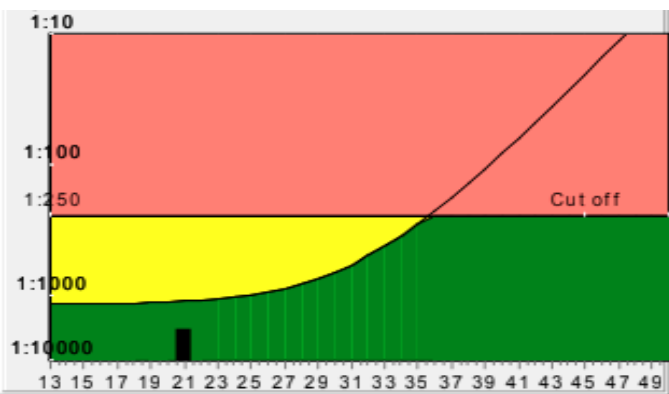
Correction factors			
Fetuses 1	IVF unknown	Previous trisomy 21 unknown	
Weight in kg 62.6	Diabetes NO	Pregnancies unknown	
Smoker NO	Origin Asian		

Biochemical Data	Ultrasound Data
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Parameter	Value	Corr Mom	
PAPP-A	4.65 mIU/ml	0.57	Gestational age 13+5
fb-hCG	23.5 ng/ml	0.96	Method CRL (<>Robinson)
			Scan date 07-01-2024

Risks at sampling date	Ultrasound Data
Age Risk 1:1121	Crown rump length in mm 77.2
Biochemical T21 risk 1:1978	Nuchal translucency MoM 0.97
Combined trisomy 21 risk 1:9095	Nasal bone PRESENT
Trisomy 13/18 + NT <1:10000	Sonographer DR.NIDHI
	Qualifications in measuring NT RADIOLOGIST

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 9095 women with the same data, there is one woman with a trisomy 21 pregnancy and 9094 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