

Date of Report 29-11-2023  
PRISCA 5.2.0.13

Patient Data			
Name	MRS AKANSHA	Patient ID	012311280114
Birthday	16-12-1983	Sample ID	11833120
Age at Sample date	40.0	Sample Date	28-11-2023
Gestational age	12+2		

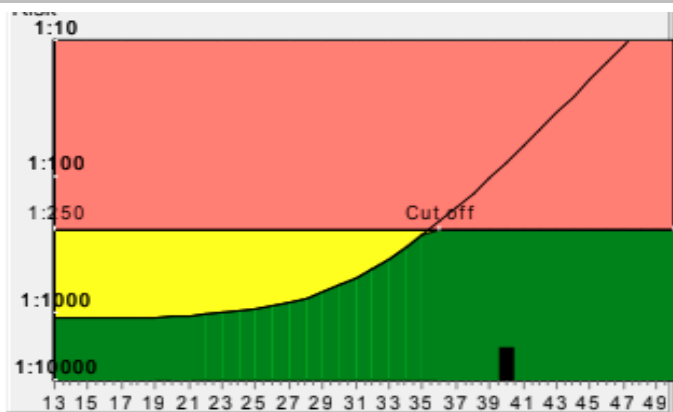
Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	92	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
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Parameter	Value	Corr Mom		
PAPP-A	4.65 mIU/ml	1.59	Gestational age	12+2
fb-hCG	26.4 ng/ml	0.75	Method	CRL (<>Robinson)
			Scan date	28-11-2023

Risks at sampling date				
Age Risk		1:79	Crown rump length in mm	58.3
Biochemical T21 risk		1:2344	Nuchal translucency MoM	1.18
Combined trisomy 21 risk		1:5356	Nasal bone	PRESENT
Trisomy 13/18 + NT		<1:10000	Sonographer	DR. VIKAS GOYAL
			Qualifications in measuring NT	MBBS

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 5356 women with the same data, there is one woman with a trisomy 21 pregnancy and 5355 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