

Date of Report 08-09-2023
PRISCA 5.2.0.13

Patient Data			
Name	MRS POOJA JINDAL	Patient ID	12309070081
Birthday	29-03-1990	Sample ID	11803895
Age at Sample date	33.4	Sample Date	07-09-2023
Gestational age	12+5		

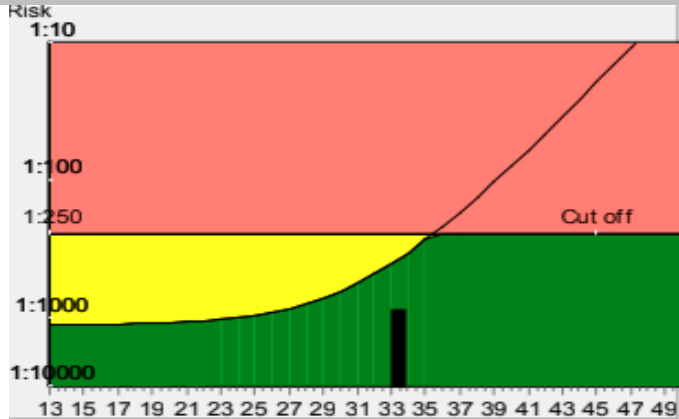
Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	57	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
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Parameter	Value	Corr Mom		
PAPP-A	2.87 mIU/ml	0.46	Gestational age	12+4
fb-hCG	55.5 ng/ml	1.54	Method	CRL (<>Robinson)
			Scan date	06-09-2023

Risks at sampling date			Ultrasound Data	
Age Risk		1:377	Crown rump length in mm	61
Biochemical T21 risk		1:134	Nuchal translucency MoM	0.63
Combined trisomy 21 risk		1:866	Nasal bone	PRESENT
Trisomy 13/18 + NT		<1:10000	Sonographer	DR.PRAKASH
			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 866 women with the same data, there is one woman with a trisomy 21 pregnancy and 865 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).

Trisomy 13/18+NT	The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values
The calculated risk for Trisomy 13/18 (with NT) is <1:10000 , which indicates a low risk	

 Risk Above Cut Off

 Risk above Age Risk

 Risk below Age risk