

Date of Report 06-09-2023
PRISCA 5.2.0.13

Patient Data			
Name	MRS RITIKA SHARMA	Patient ID	12309050223
Birth day	13-03-2001	Sample ID	11781231
Age at Sample date	22.5	Sample Date	05-09-2023
Gestational age	13+3		

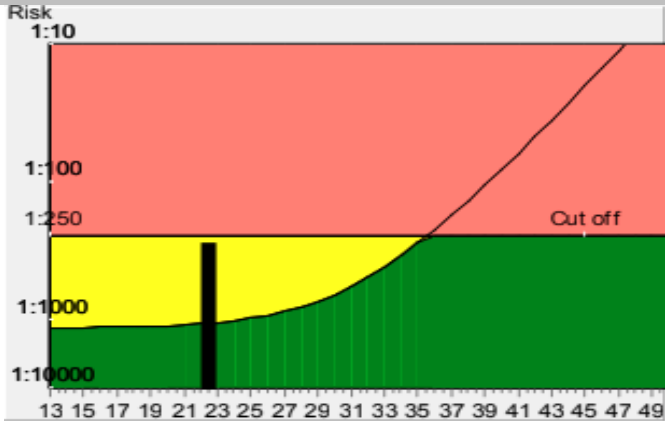
Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	42	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data	Ultrasound Data
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Parameter	Value	Corr Mom	
PAPP-A	6.38 mIU/ml	0.56	Gestational age 13+2
fb-hCG	86.2 ng/ml	2.64	Method CRL (<>Robinson)
			Scan date 04-09-2023

Risks at sampling date		
Age Risk	1:1075	Crown rump length in mm 72
Biochemical T21 risk	1:166	Nuchal translucency MoM 1.28
Combined trisomy 21 risk	1:276	Nasal bone PRESENT
Trisomy 13/18 + NT	<1:10000	Sonographer DR HARENDER
		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 276 women with the same data, there is one woman with a trisomy 21 pregnancy and 275 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 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