

Date of Report 29-10-2023
PRISCA 5.2.0.13

Patient Data			
Name	MRS KAJAL	Patient ID	012310280013
Birthday	16-06-1999	Sample ID	11843696
Age at Sample date	24.4	Sample Date	28-10-2023
Gestational age	12+3		

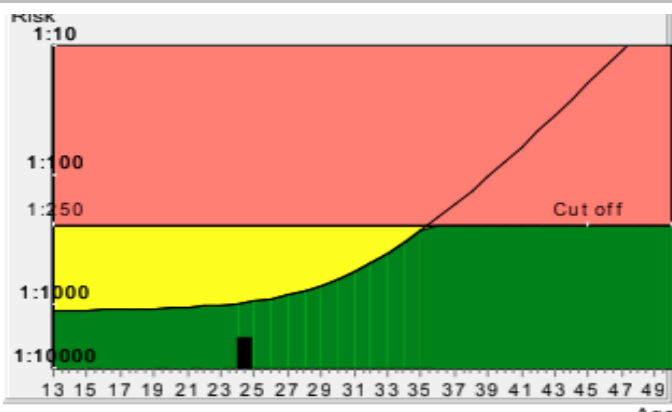
Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	49	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data		Ultrasound Data	
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Parameter	Value	Corr Mom	
PAPP-A	5.78 mIU/ml	0.87	Gestational age 12+2
fb-hCG	102.7 ng/ml	2.49	Method CRL (<>Robinson)
			Scan date 27-10-2023

Risks at sampling date		Ultrasound Data	
Age Risk	1:983	Crown rump length in mm	52.9
Biochemical T21 risk	1:494	Nuchal translucency MoM	0.71
Combined trisomy 21 risk	1:2887	Nasal bone	PRESENT
Trisomy 13/18 + NT	<1:10000	Sonographer	DR PRASHANT
		Qualifications in measuring NT	

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 2887 women with the same data, there is one woman with a trisomy 21 pregnancy and 2886 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