

Date of Report 12-10-2023
 PRISCA 5.2.0.13

Patient Data	
Name	MRS ANJALI
Birth day	10-05-2001
Age at Sample date	22.4
Gestational age	12+4

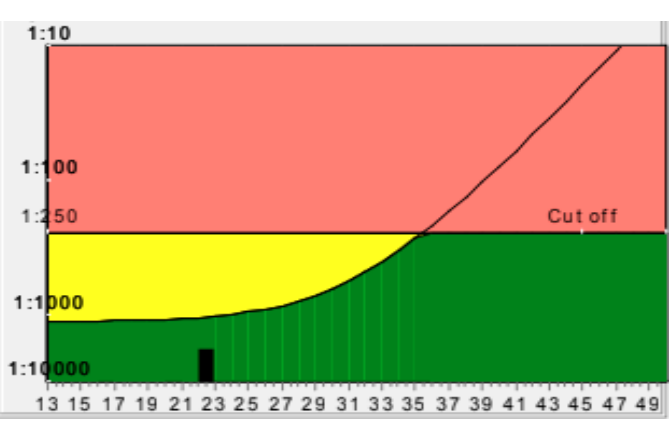
Correction factors	
Fetuses	1 IVF
Weight in kg	80 Diabetes
Smoker	NO Origin

unknown	Previous trisomy 21	unknown
NO	Pregnancies	unknown
Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+4
PAPP-A	3.84 mIU/ml	0.98	Method	CRL (<>Robinson)
fb-hCG	22.7 ng/ml	0.67	Scan date	11-10-2023

Risks at sampling date		Ultrasound Data	
Age Risk	1:1047	Crown rump length in mm	63
Biochemical T21 risk	<1:10000	Nuchal translucency MoM	1.17
Combined trisomy 21 risk	<1:10000	Nasal bone	PRESENT
Trisomy 13/18 + NT	<1:10000	Sonographer	DR SHIVANI MITTAL
		Qualifications in measuring NT	MBBS,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 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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
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Risk Above Cut Off
 Risk above Age Risk
 Risk below Age risk