

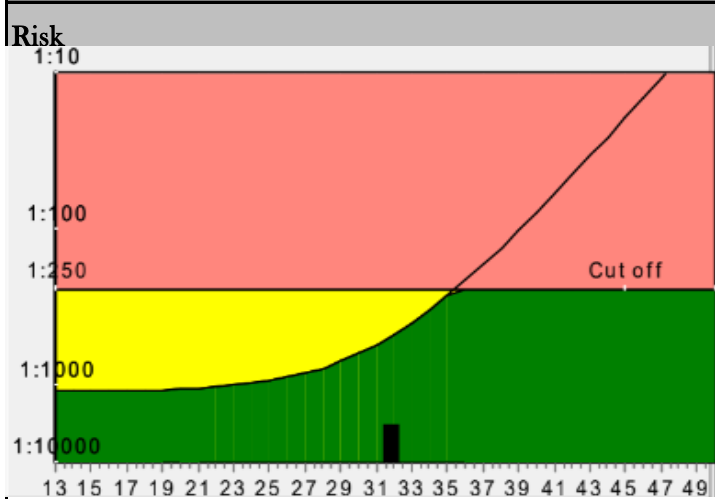
Date of Report 08-03-2024
PRISCA 5.2.0.13

Patient Data			
Name	MRS. HEMLATA	Patient ID	12403070043
Birthday	20-04-1992	Sample ID	11835454
Age at Sample date	31.9	Sample Date	07-03-2024
Gestational age	12+2		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	56.8	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	11+6
PAPP-A	4.3 mIU/ml	0.81	Method	CRL (<>Robinson)
fb-hCG	49.5 ng/ml	1.22	Scan date	04-03-2024

Risks at sampling date			
Age Risk	1:739	Crown rump length in mm	52
Biochemical T21 risk	1:1104	Nuchal translucency MoM	0.86
Combined trisomy 21 risk	1:6016	Nasal bone	PRESENT
Trisomy 13/18 + NT	<1:10000	Sonographer	DR. POONAM DHALL
		Qualifications in measuring NT	..



Trisomy 13/18+NT
The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

Down's Syndrome Risk (Trisomy 21 Screening)

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 6344 women with the same data, there is one woman with a trisomy 21 pregnancy and 6343 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 laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

