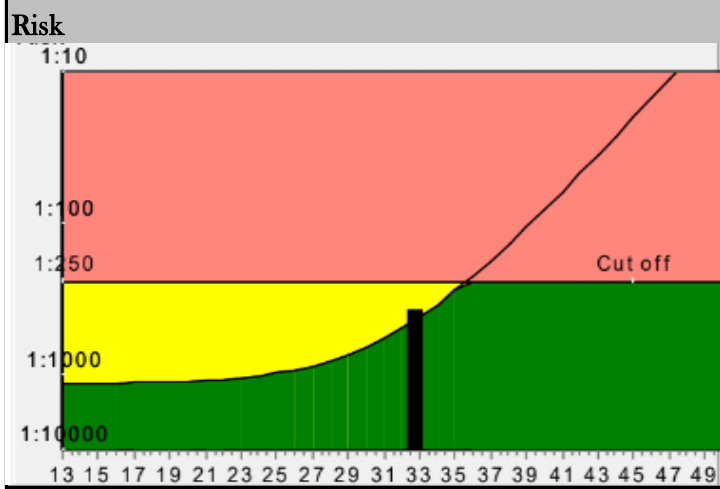


Date of Report 08-10-2024
PRISCA 5.2.0.13

Patient Data			
Name	MRS. SEEMA	Patient ID	12410070157
Birthday	05-01-1992	Sample ID	11862567
Age at Sample date	32.8	Sample Date	07-10-2024
Gestational age	13+4		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	62	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+3
PAPP-A	6.9 mIU/ml	0.91	Method	CRL (<>Robinson)
fb-hCG	123.6 ng/ml	4.57	Scan date	06-10-2024
Risks at sampling date			Crown rump length in mm	73.9
Age Risk		1:441	Nuchal translucency MoM	0.88
Biochemical T21 risk		1:68	Nasal bone	PRESENT
Combined trisomy 21 risk		1:374	Sonographer	DR.
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MBBS



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 374 women with the same data, there is one woman with a trisomy 21 pregnancy and 373 women with not affected pregnancies. The free beta HCG level is high.
 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