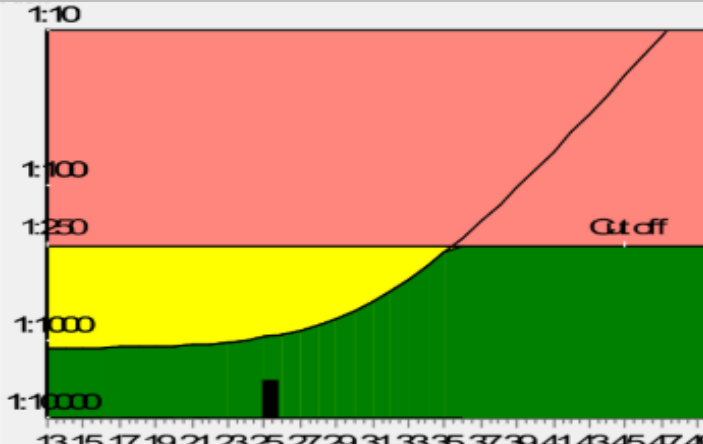


Date of Report 15-12-2024
PRISCA 5.2.0.13

Patient Data				
Name	MRS. JYOTI	Patient ID	12412140137	
Birthday	14-08-1999	Sample ID	11969943	
Age at Sample date	25.3	Sample Date	14-12-2024	
Gestational age	12+4			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	56	Diabetes	NO	Pregnancies
Smoker	NO	Origin	Asian	unknown
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+4
PAPP-A	4.3 mIU/ml	0.72	Method	CRL (<>Robinson)
fb-hCG	70.5 ng/ml	1.86	Scan date	14-12-2024
Risks at sampling date			Crown rump length in mm	62.7
Age Risk		1:948	Nuchal translucency MoM	1.05
Biochemical T21 risk		1:637	Nasal bone	PRESENT
Combined trisomy 21 risk		1:2338	Sonographer	DR. DEEPAK
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD
Risk			Down's Syndrome Risk (Trisomy 21 Screening)	
			<p>The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 2338 women with the same data, there is one woman with a trisomy 21 pregnancy and 2337 women with not affected pregnancies.</p> <p>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!</p> <p>The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).</p>	
Trisomy 13/18+NT			The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values	
The calculated risk for Trisomy 13/18 (with NT) is <1:10000 , which indicates a low risk				



Risk Above Cut Off



Risk above Age Risk



Risk below Age risk