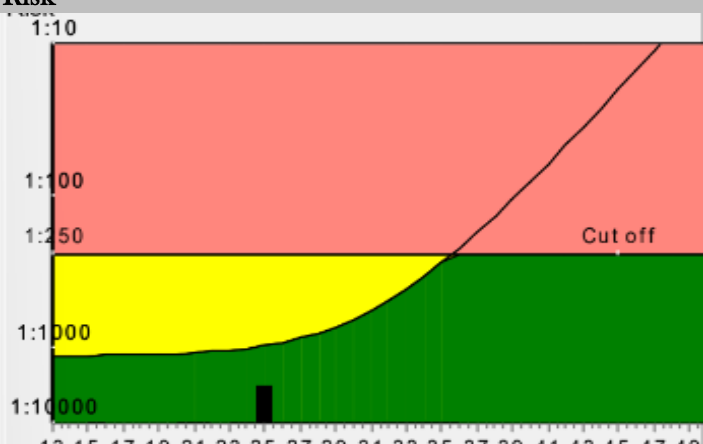


Date of Report 11-05-2024
PRISCA 5.2.0.13

Patient Data					
Name	MRS. PRIYANKA SINGH		Patient ID	012405100056	
Birthday	15-06-199		Sample ID	012001331	
Age at Sample date	24.9		Sample Date	10-05-2024	
Gestational age	13+3				
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	50	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	13+1	
PAPP-A	6.5 mIU/ml	0.70	Method	CRL (<>Robinson)	
fb-hCG	52.7 ng/ml	1.72	Scan date	08-05-2024	
Risks at sampling date			Crown curm length in mm	70.4	
Age Risk	1:995		Nuchal translucency MoM	0.74	
Bicochemical T21 risk	1:764		Nasal bone	present	
Commbined Trisomy 21 risk	1:4505		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 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 4505 women with the same data, there is one woman with a trisomy 21 pregnancy and 4504 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