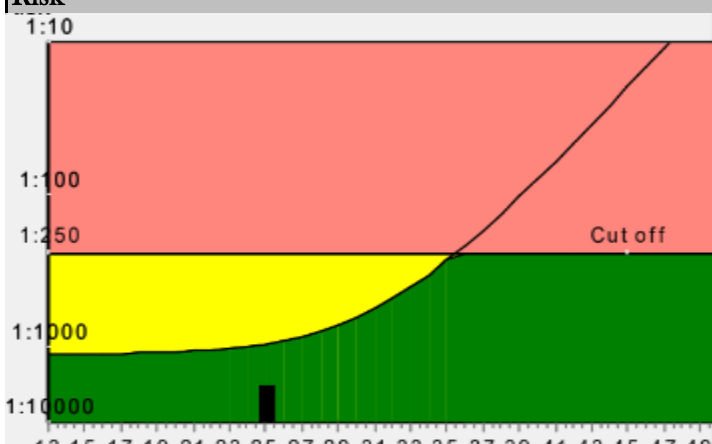


Date of Report 13-04-2024
PRISCA 5.2.0.13

Patient Data					
Name	MRS. NISHA		Patient ID	012404110215	
Birthday	26-03-1999		Sample ID	11848135	
Age at Sample date	25.0		Sample Date	11-04-2024	
Gestational age	12+5				
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	50.8	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	12+4	
PAPP-A	4.8 mIU/ml	0.38	Method	CRL (<>Robinson)	
fb-hCG	53.9 ng/ml	1.43	Scan date	10-04-2024	
Risks at sampling date			Crown rump length in mm	62	
Age Risk	1:966		Nuchal translucency MoM	0.69	
Biochemical T21 risk	1:1074		Nasal bone	PRESENT	
Combined trisomy 21 risk	1:6222		Sonographer	DR.RAJENDER KUMAR	
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	MBBS	
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 6222 women with the same data, there is one woman with a trisomy 21 pregnancy and 6221 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