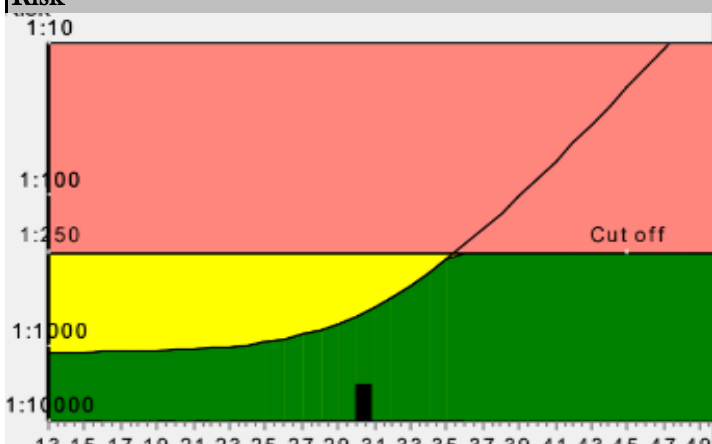


Date of Report 07-05-2024
PRISCA 5.2.0.13

Patient Data					
Name	MRS. NANCY DHAWAN F1		Patient ID	012405060084	
Birthday	08-12-1993		Sample ID	11876965	
Age at Sample date	30.4		Sample Date	06-05-2024	
Gestational age	12+3				
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	62	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	12+2	
PAPP-A	5.1 mIU/ml	0.54	Method	CRL (<>Robinson)	
fb-hCG	63.4 ng/ml	0.77	Scan date	05-05-2024	
Risks at sampling date			Crown curm length in mm	58	
Age Risk	1:611		Nuchal translucency MoM	0.79	
Bicochemical T21 risk	1:1570		Nasal bone	present	
Commbined Trisomy 21 risk	1:8830		Sonographer	DR.INDRAJEET	
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 8830 women with the same data, there is one woman with a trisomy 21 pregnancy and 8829 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