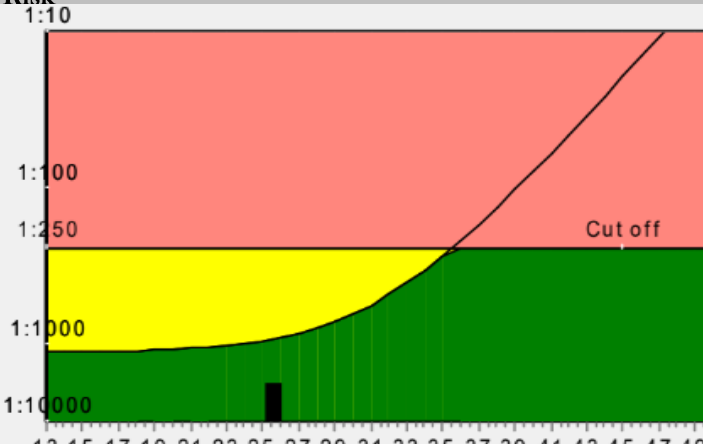


Date of Report 01-03-2024  
PRISCA 5.2.0.13

Patient Data					
Name	<b>MRS. PRIYA KUMARI</b>		Patient ID	12402280132	
Birthday	15-08-1998		Sample ID	11850944	
Age at Sample date	25.5		Sample Date	28-02-2024	
Gestational age	12+6				
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	56.5	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	12+5	
PAPP-A	5.3 mIU/ml	0.80	Method	CRL (<>Robinson)	
fb-hCG	59.6 ng/ml	1.71	Scan date	28-02-2024	
Risks at sampling date			Crown rump lenth in mm	63.3	
Age Risk	1:948		Nuchal translucency MoM	0.93	
Biochemical T21 risk	1:1023		Nasal bone	present	
Combined trisomy T21 risk	1:4961		Sonographer	DR. PRAVEEN	
Trisomy 13/18 + NT	<1:10000		Qualification in measuring NT	C/R	
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
			<p><b>The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 4961 women with the same data, there is one woman with a trisomy 21 pregnancy and 4960 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 the NT measurement was done according to accepted guidelines (prenat Diagn 18 : 511 - 523 (1998)). The laboratory can not be hold responsible for their impact on risk assessment ! Calculated risk have no diagnostic value!</p>		
Trisomy 13/18 + NT					
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