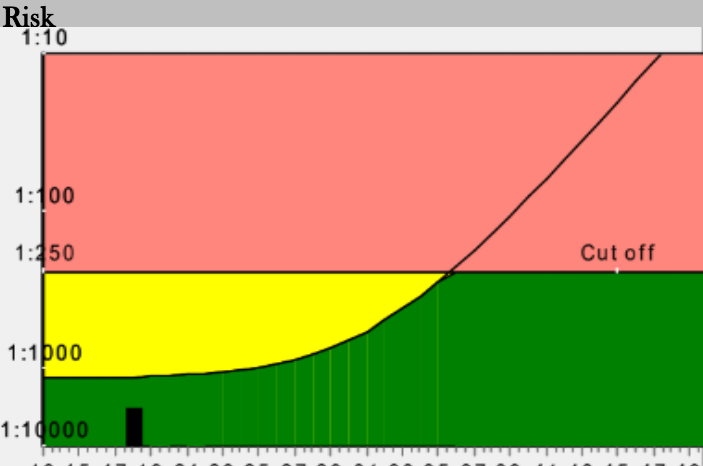


Date of Report 12-03-2024  
PRISCA 5.2.0.13

Patient Data					
Name	MRS. KAVITA	Patient ID	12403090176		
Birthday	04-02-2006	Sample ID	11823478		
Age at Sample date	18.1	Sample Date	09-03-2024		
Gestational age	13+6				
Correction factors					
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown	
Weight in kg	52	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	13+5	
PAPP-A	5.8 mIU/ml	0.57	Method	CRL (<>Robinson)	
fb-hCG	19.5 ng/ml	0.75	Scan date	09-03-2024	
Risks at sampling date			Crown rump length in mm	77.1	
Age Risk	1:1156		Nuchal translucency MoM	0.59	
Biochemical T21 risk	1:3524		Nasal bone	present	
combined trisomy 21 risk	<1:10000		Sonographer	DR.	
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 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</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