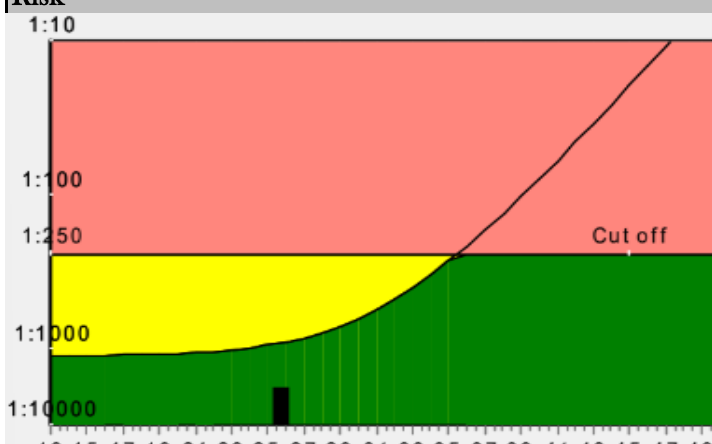


Date of Report 29-03-2024  
PRISCA 5.2.0.13

Patient Data					
Name	MRS. POOJA	Patient ID	12403270309		
Birthday	08-07-1998	Sample ID	11818084		
Age at Sample date	25.7	Sample Date	27-03-2024		
Gestational age	12+4				
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	12+3	
PAPP-A	5.3 mIU/ml	0.77	Method	CRL (<>Robinson)	
fb-hCG	42.9 ng/ml	1.09	Scan date	27-03-2024	
Risks at sampling date			Crown rump length in mm	59.7	
Age Risk	1:930		Nuchal translucency MoM	1.09	
Biochemical T21 risk	1:2688		Nasal bone	PRESENT	
Combined trisomy 21 risk	1:8622		Sonographer	DR. VIKASH	
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 (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 8622 women with the same data, there is one woman with a trisomy 21 pregnancy and 8621 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