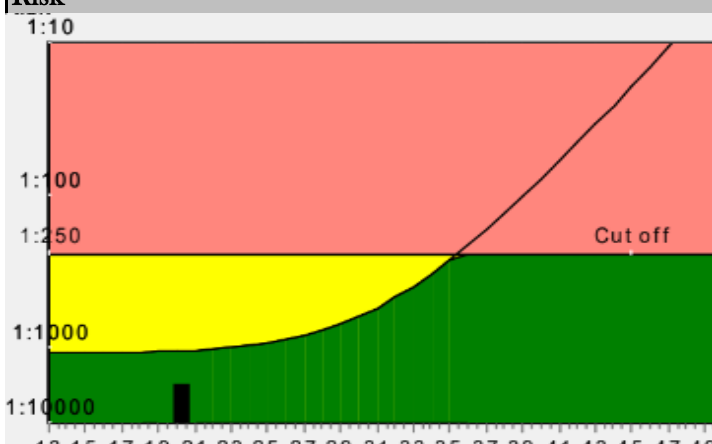


Date of Report 14-04-2024
PRISCA 5.2.0.13

Patient Data					
Name	MRS. ANCHAL	Patient ID	12404120257		
Birthday	01-01-2004	Sample ID	12001676		
Age at Sample date	20.3	Sample Date	12-04-2024		
Gestational age	12+1				
Correction factors					
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown	
Weight in kg	43	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	11+3	
PAPP-A	5.1 mIU/ml	0.74	Method	CRL (<>Robinson)	
fb-hCG	128.5 ng/ml	2.76	Scan date	08-04-2024	
Risks at sampling date			Crown rump length in mm	48.8	
Age Risk		1:1070	Nuchal translucency MoM	0.68	
Biochemical T21 risk		1:288	Nasal bone	PRESENT	
Combined trisomy 21 risk		1:1779	Sonographer	DR. DEEPIKA	
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 1779 women with the same data, there is one woman with a trisomy 21 pregnancy and 1778 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