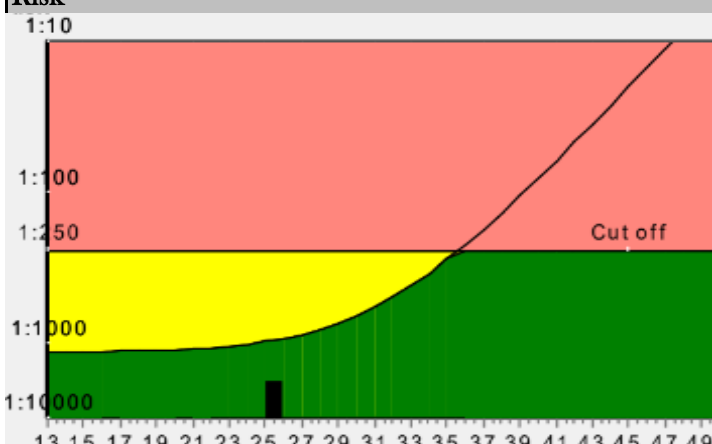


Date of Report 04-04-2024  
PRISCA 5.2.0.13

Patient Data					
Name	<b>MRS. SHRADHA</b>		Patient ID	12404020244	
Birthday	04-10-1998		Sample ID	11816077	
Age at Sample date	25.5		Sample Date	02-04-2024	
Gestational age	13+4				
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	64	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		
Biochemical Data			Ultrasound Data		
<b>Parameter</b>	<b>Value</b>	<b>Corr Mom</b>	Gestational age	13+4	
PAPP-A	6.9 mIU/ml	0.94	Method	CRL (<>Robinson)	
fb-hCG	43.5 ng/ml	1.62	Scan date	02-04-2024	
Risks at sampling date			Crown rump length in mm	74	
Age Risk	1:972		Nuchal translucency MoM	0.73	
Biochemical T21 risk	1:1720		Nasal bone	present	
Combined trisomy 21 risk	1:9299		Sonographer	DR. INDERJEET	
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	MD	
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 9299 women with the same data, there is one woman with a trisomy 21 pregnancy and 9298 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		
<p>The calculated risk for Trisomy 13/18 (with NT) is &lt;1:10000 , which indicates a low risk</p>					

