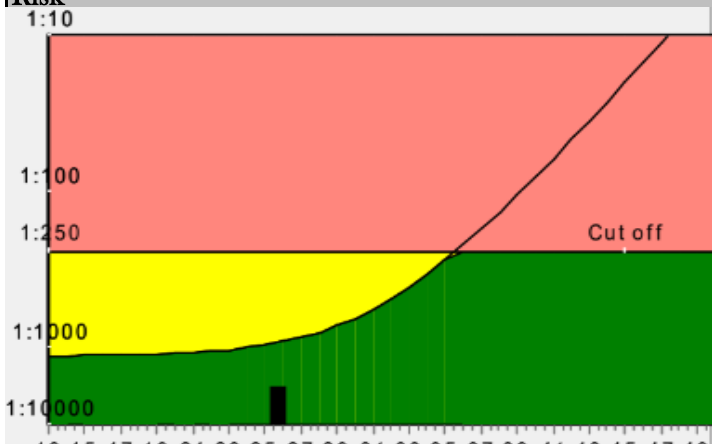


Date of Report 19-03-2024
PRISCA 5.2.0.13

Patient Data				
Name	MRS. SHILPI NAMDEV		Patient ID	12403170207
Birthday	16-07-1998		Sample ID	11832340
Age at Sample date	25.7		Sample Date	17-03-2024
Gestational age	13+2			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	47	Diabetes	NO	Pregnancies
Smoker	NO	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+2
PAPP-A	6.1 mIU/ml	0.64	Method	CRL (<>Robinson)
fb-hCG	28.6 ng/ml	0.87	Scan date	17-03-2024
Risks at sampling date			Crown rump length in mm	71.3
Age Risk	1:955		Nuchal translucency MoM	0.73
Biochemical T21 risk	1:2867		Nasal bone	PRESENT
Combined trisomy 21 risk	<1:10000		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 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