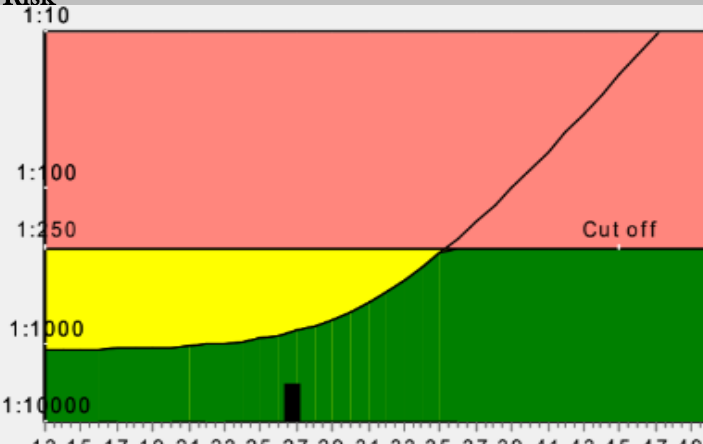


Date of Report 03-03-2024  
PRISCA 5.2.0.13

Patient Data				
Name	<b>MRS. BHAWNA SHARMA</b>		Patient ID	12403020198
Birthdate	02-06-1997		Sample ID	11818917
Age at Sample date	26.7		Sample Date	02-03-2024
Gestational age	11+5			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	58	Diabetes	NO	Pregnancies
Smoker	NO	Origin	Asian	unknown
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	11+5
PAPP-A	3.8 mIU/ml	0.93	Method	CRL (<>Robinson)
fb-hCG	57.6 ng/ml	1.24	Scan date	02-03-2024
Risks at sampling date			Crown rump length in mm	49
Age Risk	1:848		Nuchal translucency MoM	0.64
Biochemical T21 risk	1:2764		Nasal bone	PRESENT
Combined trisomy 21 risk	<1:10000		Sonographer	DR. SAGAR
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	MBBS,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 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	
<p>The calculated risk for Trisomy 13/18 (with NT) is &lt;1:10000 , which indicates a low risk</p>				

Risk Above Cut Off
  Risk above Age Risk
  Risk below Age risk