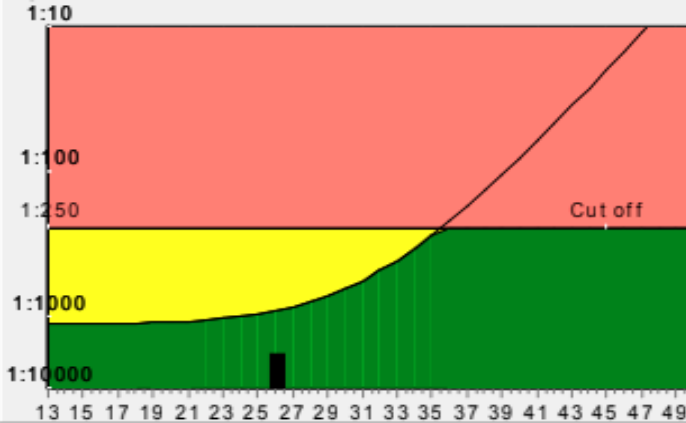


Date of Report 7/1/2024
PRISCA 5.2.0.13

Patient Data				
Name	BHAWNA	Patient ID	12401060087	
Birth day	3/12/1997	Sample ID	11827715	
Age at sample	26.1	Sample Date	6/1/2024	
Gestational age	13+0			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	83 Diabetes	NO	Pregnancies	unknown
Smoker	NO Origin	Asian		
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+6
PAPP-A	4.12 mIU/ml	0.94	Method	CRL (<>Robinson)
fb-hCG	35.9 ng/ml	1.21	Scan date	5/1/2024
Risks at sampling date			Crown rump length in mm	64.9
Age Risk	1:925		Nuchal translucency MoM	1.08
Biochemical T21 risk	1:171		Nasal bone	Present
Combined trisomy 21 risk	<1:10000		Sonographer	DRJYOTI
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	MBBS
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 more than 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> <p>The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values</p>	
Trisomy 13/18 + NT				
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