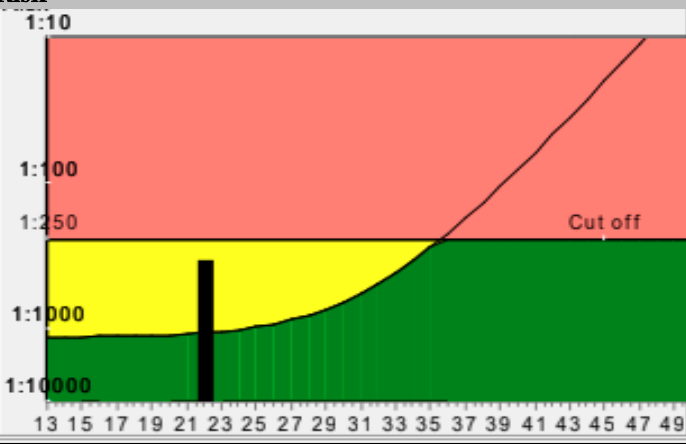


Date of Report 5/5/2024
PRISCA 5.2.0.13

Patient Data				
Name	JYOTI	Patient ID	012405030229	
Birthday	1/4/2002	Sample ID	12001852	
Age at sample	22.1	Sample Date	3/5/2024	
Gestational age	13+3			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	53	Diabetes	NO	Pregnancies
Smoker	NO	Origin	Asian	unknown
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+1
PAPP-A	7.50 mIU/ml	0.86	Method	CRL (<>Robinson)
fb-hCG	113.5 ng/ml	3.79	Scan date	24/04/2024
Risks at sampling date			Crown rump length in mm	56
Age Risk	1:1084		Nuchal translucency MoM	1.22
Biochemical T21 risk	1:172		Nasal bone	Present
Combined trisomy 21 risk	1:342		Sonographer	DR. HARENDRA BHASKAR
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 342 women with the same data, there is one woman with a trisomy 21 pregnancy and 341 women with not affected pregnancies. The free beta HCG level is high. 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