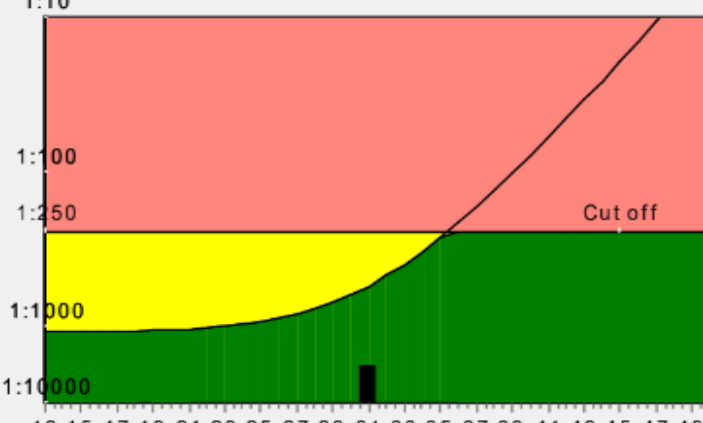


Date of Report 15-04-2024  
PRISCA 5.2.0.13

Patient Data				
Name	<b>MRS. MAITREYEE KULKARNI</b>		Patient ID	012404140076
Birthday	0-05-1993		Sample ID	11835067
Age at Sample date	31.0		Sample Date	14-04-2024
Gestational age	12+1			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	67	Diabetes	NO	Pregnancies
Smoker	NO	Origin	Asian	unknown
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+0
PAPP-A	4.5 mIU/ml	1.10	Method	CRL (<>Robinson)
fb-hCG	18.5 ng/ml	0.46	Scan date	13-04-2024
Risks at sampling date			Crown rump length in mm	54
Age Risk		1:561	Nuchal translucency MoM	0.77
Biochemical T21 risk		<1:10000	Nasal bone	PRESENT
Combined trisomy 21 risk		<1:10000	Sonographer	DR. DHRUV TANEJA
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 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