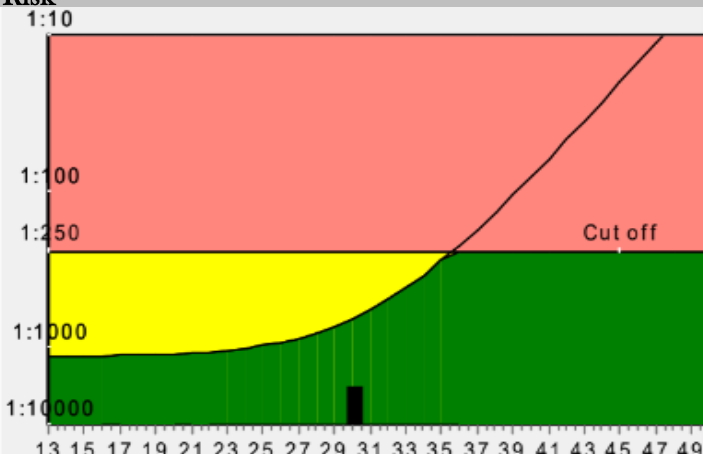


Date of Report 12-03-2024
PRISCA 5.2.0.13

Patient Data				
Name	MRS. PALLAVI KUMARI		Patient ID	12403100153
Birthday	12-01-1994		Sample ID	11823196
Age at Sample date	30.2		Sample Date	10-03-2024
Gestational age	13+4			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21 unknown
Weight in kg	57	Diabetes	NO	Pregnancies unknown
Smoker	NO	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+3
PAPP-A	6.8 mIU/ml	0.81	Method	CRL (<>Robinson)
fb-hCG	14.6 ng/ml	0.52	Scan date	09-03-2024
Risks at sampling date			Crown rump length in mm	74
Age Risk	1:656		Nuchal translucency MoM	0.87
Biochemical T21 risk	<1:10000		Nasal bone	present
combined trisomy 21 risk	<1:10000		Sonographer	DR. HARENDERA 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 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 <1:10000, which indicates a low risk</p>				

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