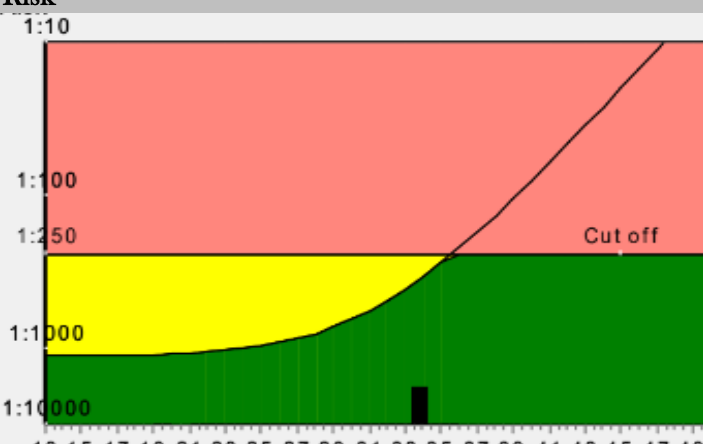


Date of Report 10-05-2024
PRISCA 5.2.0.13

Patient Data				
Name	MRS. ANJU DEVI		Patient ID	12405090201
Birthday	20-08-1990		Sample ID	11888373
Age at Sample date	33.7		Sample Date	09-05-2024
Gestational age	13+1			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21 unknown
Weight in kg	73	Diabetes	NO	Pregnancies unknown
Smoker	NO	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+0
PAPP-A	6.1 mIU/ml	1.13	Method	CRL (<>Robinson)
fb-hCG	31.8 ng/ml	1.08	Scan date	01-05-2024
Risks at sampling date			Crown curm length in mm	52
Age Risk	1:363		Nuchal translucency MoM	1.22
Bicochemical T21 risk	1:2484		Nasal bone	present
Commbined Trisomy 21 risk	1:5087		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 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 5087 women with the same data, there is one woman with a trisomy 21 pregnancy and 5086 women with not affected pregnancies.</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