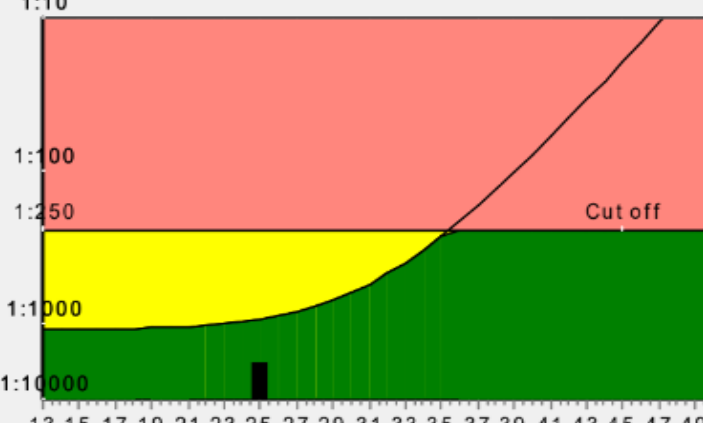


Date of Report 31-03-2024
PRISCA 5.2.0.13

Patient Data				
Name	MRS. N. PALLAVI		Patient ID	12403300043
Birthday	27-04-1999		Sample ID	11494374
Age at Sample date	24.9		Sample Date	30-03-2024
Gestational age	12+1			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21 unknown
Weight in kg	45	Diabetes	NO	Pregnancies unknown
Smoker	NO	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+0
PAPP-A	5.1 mIU/ml	0.78	Method	CRL (<>Robinson)
fb-hCG	42.1 ng/ml	0.92	Scan date	29-03-2024
Risks at sampling date			Crown rump length in mm	53.2
Age Risk	1:951		Nuchal translucency MoM	0.49
Biochemical T21 risk	1:4119		Nasal bone	PRESENT
Combined trisomy 21 risk	<1:10000		Sonographer	DR.
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