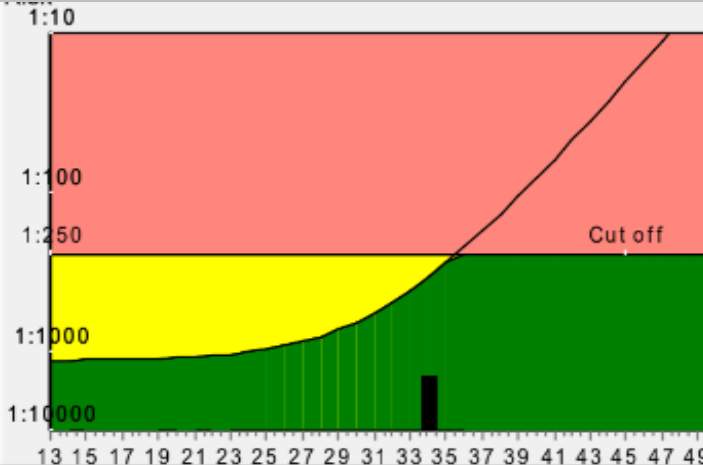


Date of Report 15-07-2024
PRISCA 5.2.0.13

Patient Data				
Name	MRS. MANJUBALA		Patient ID	12407140076
Birthday	28-06-1990		Sample ID	11787446
Age at Sample date	34.0		Sample Date	14-07-2024
Gestational age	13+2			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	57	Diabetes	NO	Pregnancies
Smoker	NO	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+1
PAPP-A	6.95 mIU/ml	0.91	Method	CRL(<>Robinson)
fb-hCG	68.4 ng/ml	2.24	Scan date	13-07-2024
Risks at sampling date			Crown rump length in mm	68.8
Age Risk		1:342	Nuchal translucency MoM	0.58
Biochemical T21 risk		1:251	Nasal bone	present
Combined trisomy 21 risk		1:1428	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 test (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 1428 women with the same data, there is one woman with a trisomy 21 pregnancy and 1427 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 held 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