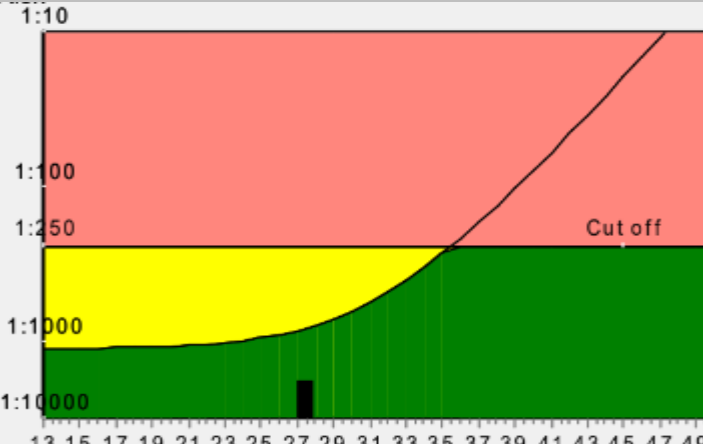


Date of Report 24-06-2024  
PRISCA 5.2.0.13

Patient Data				
Name	MRS. PARUL	Patient ID	12406230104	
Birthday	01-01-1997	Sample ID	12002071	
Age at Sample date	27.5	Sample Date	23-06-2024	
Gestational age	12+4			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	46	Diabetes	NO	Pregnancies
Smoker	NO	Origin	Asian	unknown
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+4
PAPP-A	5.3 mIU/ml	0.70	Method	CRL (<>Robinson)
fb-hCG	55.9 ng/ml	1.38	Scan date	23-06-2024
Risks at sampling date			Crown rump length in mm	59.6
Age Risk		1:831	Nuchal translucency MoM	0.64
Biochemical T21 risk		1:1119	Nasal bone	PRESENT
Combined trisomy 21 risk		1:6377	Sonographer	DR. Rohit Aggarwal
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD
Risk			Down's Syndrome Risk (Trisomy 21 Screening)	
			<p><b>The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 6377 women with the same data, there is one woman with a trisomy 21 pregnancy and 6376 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