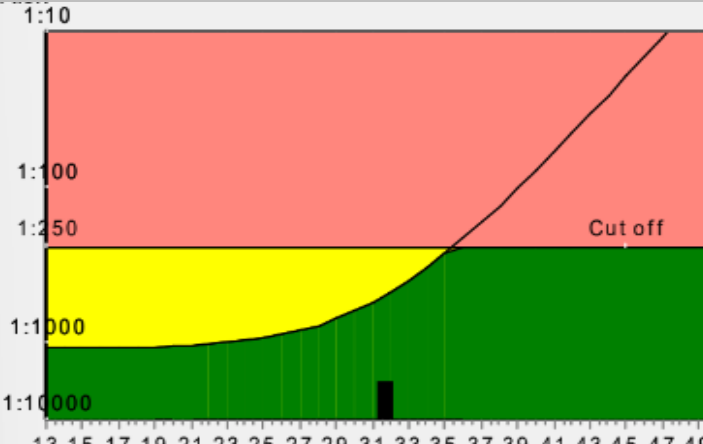


Date of Report 21-09-2024  
PRISCA 5.2.0.13

Patient Data					
Name	MRS. MAMTA	Patient ID	12409190100		
Birthday	01-01-1993	Sample ID	11860358		
Age at Sample date	31.7	Sample Date	19-09-2024		
Gestational age	12+2				
Correction factors					
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown	
Weight in kg	63	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	12+1	
PAPP-A	4.8 mIU/ml	1.03	Method	CRL (<>Robinson)	
fb-hCG	54.3 ng/ml	1.39	Scan date	18-09-2024	
Risks at sampling date			Crown rump length in mm	57	
Age Risk		1:503	Nuchal translucency MoM	0.89	
Biochemical T21 risk		1:1556	Nasal bone	PRESENT	
Combined trisomy 21 risk		1:7528	Sonographer	DR. ARSHIBIR SINGH	
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 more 7528 women with the same data, there is one woman with a trisomy 21 pregnancy and 7527 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