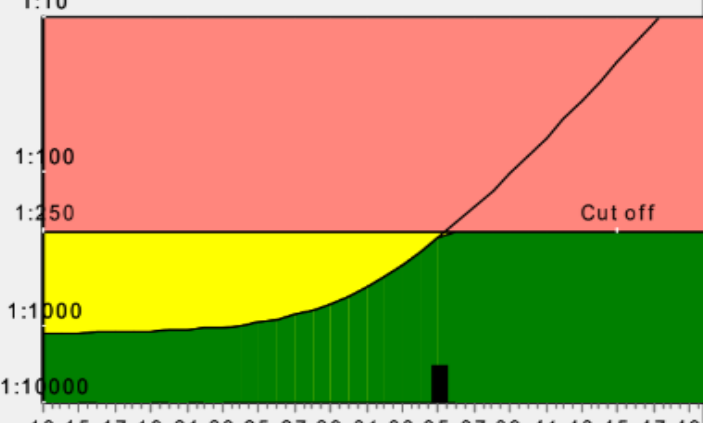


Date of Report 14-03-2024  
PRISCA 5.2.0.13

Patient Data					
Name	MRS. POONAM		Patient ID	12403130134	
Birthday	15-02-1989		Sample ID	11812538	
Age at Sample date	35.1		Sample Date	13-03-2024	
Gestational age	12+3				
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	77	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	12+2	
PAPP-A	5.3 mIU/ml	1.37	Method	CRL (<>Robinson)	
fb-hCG	23.5 ng/ml	0.66	Scan date	12-03-2024	
Risks at sampling date			Crown rump length in mm	58	
Age Risk	1:267		Nuchal translucency MoM	0.92	
Biochemical T21 risk	1:7750		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