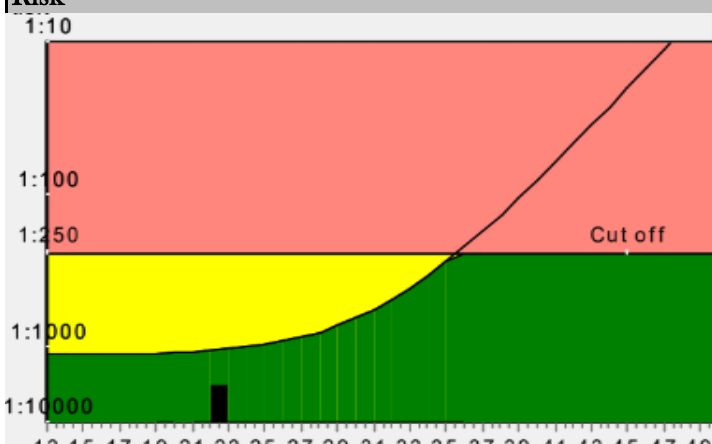


Date of Report 21-03-2024  
PRISCA 5.2.0.13

Patient Data				
Name	MRS. BHAWNA	Patient ID	12403200166	
Birthday	24-09-2001	Sample ID	11823159	
Age at Sample date	22.5	Sample Date	20-03-2024	
Gestational age	13+1			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	64 Diabetes	NO	Pregnancies	unknown
Smoker	NO Origin	Asian		
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+1
PAPP-A	5.6 mIU/ml	0.89	Method	CRL (<>Robinson)
fb-hCG	61.5 ng/ml	2	Scan date	20-03-2024
Risks at sampling date			Crown rump length in mm	67.8
Age Risk		1:1065	Nasal translucency MoM	0.87
Biochemical T21 risk		1:976	Nasal bone	present
Combined trisomy 21 risk		1:5121	Sonographer	DR. RAVIKANT
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	MD
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 it is expected that among 5121 women with the same data, there is one woman with a trisomy 21 pregnancy and 5120 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 the NT measurement was done according to accepted guidelines (prenat Diagn 18:511-523(1998)). The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>	
Trisomy 13/18 + NT				
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