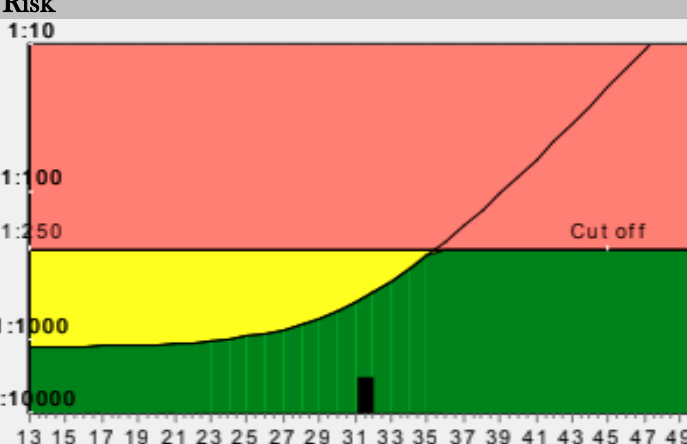


Date of Report 28/09/2023
PRISCA 5.2.0.13

Patient Data				
Name	MAMTA YADAV	Patient ID	012309270176	
Birthday	23/03/1992	Sample ID	11787478	
Age at sample	31.5	Sample Date	27/09/2023	
Gestational age	12+4			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	49 Diabetes	NO	Pregnancies	unknown
Smoker	NO Origin	Asian		
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+4
PAPP-A	3.92 mIU/ml	0.56	Method	CRL (<>Robinson)
fb-hCG	36.4 ng/ml	0.92	Scan date	27/09/2023
Risks at sampling date			Crown rump length in mm	61.8
Age Risk	1:524		Nuchal translucency MoM	0.88
Biochemical T21 risk	1:998		Nasal bone	Present
Combined trisomy 21 risk	1:5391		Sonographer	DR VIKAS GOYAL
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 5391 women with the same data, there is one woman with a trisomy 21 pregnancy and 5390 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> <p>The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values</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