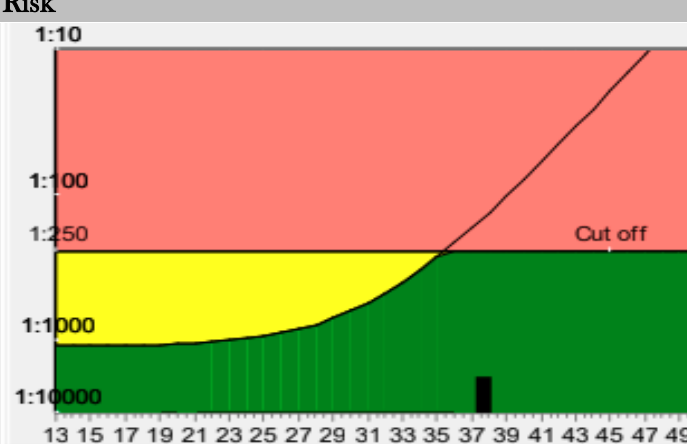


Date of Report 5/2/2024
PRISCA 5.2.0.13

Patient Data				
Name	URMILA	Patient ID	012402020126	
Birthday	21/05/1986	Sample ID	11818717	
Age at sample	37.7	Sample Date	2/3/2024	
Gestational age	12+2			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	86.3 Diabetes	NO	Pregnancies	unknown
Smoker	NO Origin	Asian		
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+6
PAPP-A	4.50 mIU/ml	1.42	Method	CRL (<>Robinson)
fb-hCG	51.7 ng/ml	1.44	Scan date	16/12/2023
Risks at sampling date			Crown rump length in mm	70
Age Risk	1:142		Nuchal translucency MoM	0.91
Biochemical T21 risk	1:761		Nasal bone	Present
Combined trisomy 21 risk	1:3656		Sonographer	DR SANJEEV KUMAR
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 3656 women with the same data, there is one woman with a trisomy 21 pregnancy and 3655 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>	
			<p>Trisomy 13/18 + NT</p> <p>The calculated risk for Trisomy 13/18 (with NT) is < 1:10000, which indicates a low risk</p>	

 Risk Above Cut Off

 Risk above Age Risk

 Risk below Age risk