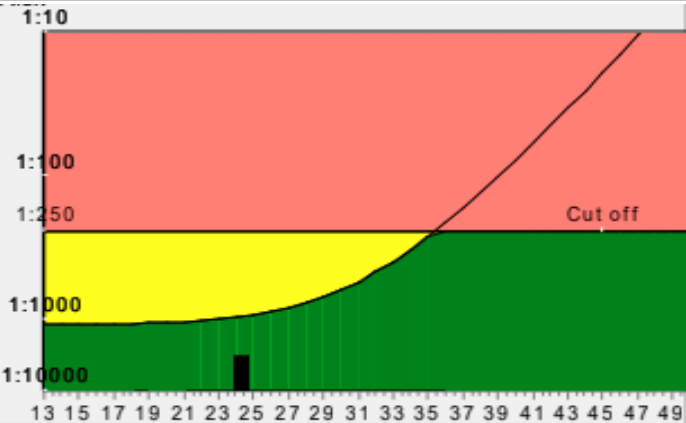


Date of Report 13/10/2023
PRISCA 5.2.0.13

Patient Data				
Name	PRIYA KUMARI	Patient ID	012310090176	
Birthday	13/07/1999	Sample ID	11836822	
Age at sample	24.2	Sample Date	9/10/2023	
Gestational age	12+1			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	54.7	Diabetes	NO	Pregnancies
Smoker	NO	Origin	Asian	unknown
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+1
PAPP-A	3.80 mIU/ml	0.73	Method	CRL (<>Robinson)
fb-hCG	41.5 ng/ml	0.97	Scan date	9/10/2023
Risks at sampling date			Crown rump length in mm	55.8
Age Risk	1:978		Nuchal translucency MoM	0.57
Biochemical T21 risk	1:3152		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 more than 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> <p>The laboratory cannot be held 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