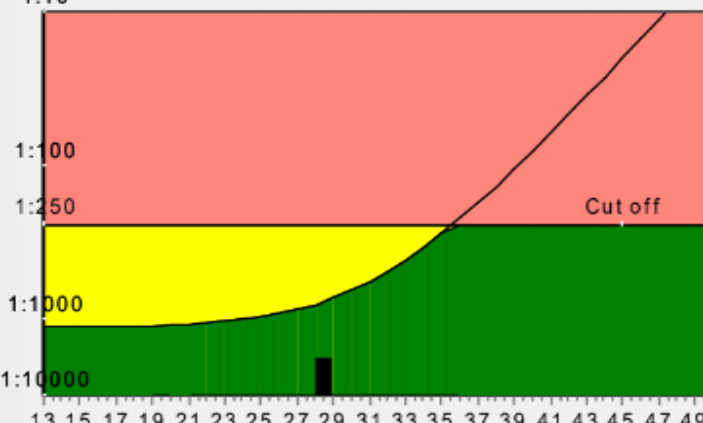


Date of Report 12-09-2024
PRISCA 5.2.0.13

Patient Data				
Name	MRS. POOJA KUMARI		Patient ID	12409110243
Birthday	05-04-1996		Sample ID	11966974
Age at Sample date	28.4		Sample Date	11-09-2024
Gestational age	13+1			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21 unknown
Weight in kg	62.3	Diabetes	NO	Pregnancies unknown
Smoker	NO	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+1
PAPP-A	5.4 mIU/ml	0.83	Method	CRL (<>Robinson)
fb-hCG	61.84 ng/ml	2	Scan date	11-09-2024
Risks at sampling date			Crown rump length in mm	69.3
Age Risk		1:781	Nuchal translucency MoM	0.86
Biochemical T21 risk		1:619	Nasal bone	PRESENT
Combined trisomy 21 risk		1:3361	Sonographer	DR. AMENDA
Trisomy 13/18 + NT		<1:10000	Qualifications 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 (with NT) it is expected that among 3361 women with the same data, there is one woman with a trisomy 21 pregnancy and 3360 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>	
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