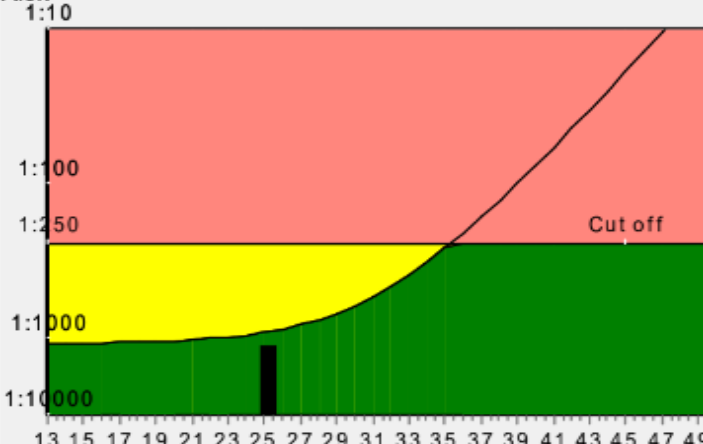


Date of Report 16-06-2024  
PRISCA 5.2.0.13

Patient Data				
Name	MRS. SHILPI SARKAR		Patient ID	12406150133
Birthday	22-04-1999		Sample ID	11870797
Age at Sample date	25.1		Sample Date	15-06-2024
Gestational age	11+5			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	42	Diabetes	NO	Pregnancies
Smoker	NO	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	11+5
PAPP-A	4.8 mIU/ml	0.82	Method	CRL (<>Robinson)
fb-hCG	108 ng/ml	2.07	Scan date	15-06-2024
Risks at sampling date			Crown rump length in mm	50
Age Risk		1:927	Nuchal translucency MoM	1.26
Biochemical T21 risk		1:648	Nasal bone	PRESENT
Combined trisomy 21 risk		1:1150	Sonographer	DR.
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MBBS
Risk			Down's Syndrome Risk (Trisomy 21 Screening)	
			<p><b>The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 1150 women with the same data, there is one woman with a trisomy 21 pregnancy and 1149 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