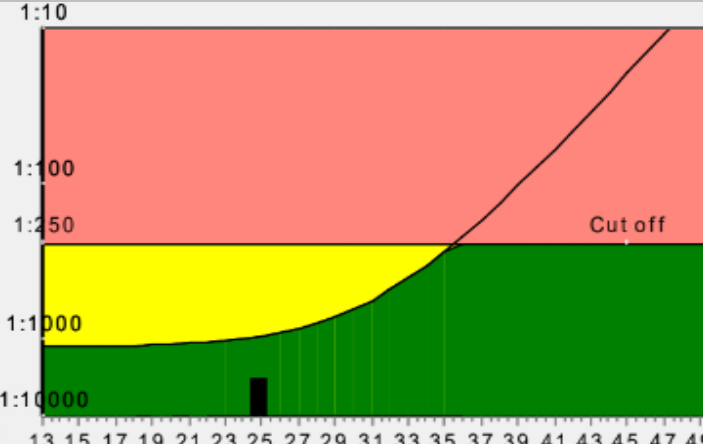


Date of Report 21-09-2024
PRISCA 5.2.0.13

Patient Data					
Name	MRS. PAYAL	Patient ID	12409180187		
Birthday	07-11-1999	Sample ID	11978947		
Age at Sample date	24.9	Sample Date	18-09-2024		
Gestational age	12+6				
Correction factors					
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown	
Weight in kg	70	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	12+6	
PAPP-A	5.9 mIU/ml	1.15	Method	CRL (<>Robinson)	
fb-hCG	38.6 ng/ml	1.19	Scan date	18-09-2024	
Risks at sampling date			Crown rump length in mm	65.1	
Age Risk		1:978	Nuchal translucency MoM	0.74	
Biochemical T21 risk		1:5539	Nasal bone	PRESENT	
Combined trisomy 21 risk		<1:10000	Sonographer	DR. PARUL	
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>		
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