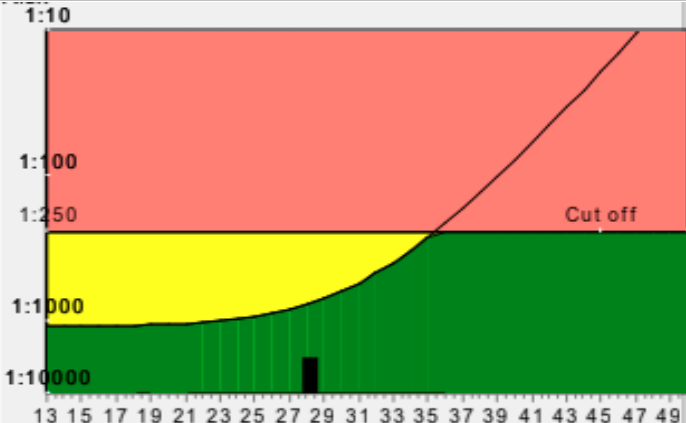


Date of Report 16/1/2024  
PRISCA 5.2.0.13

Patient Data				
Name	ARPITA	Patient ID	12401150022	
Birthday	19/10/1995	Sample ID	11827479	
Age at sample	28.20	Sample Date	15/1/2024	
Gestational age	12+1			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	52.7	Diabetes	NO	Pregnancies
Smoker	NO	Origin	Asian	unknown
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+0
PAPP-A	3.90 mIU/ml	0.72	Method	CRL (<>Robinson)
fb-hCG	41.2 ng/ml	0.95	Scan date	14/1/2024
Risks at sampling date			Crown rump length in mm	54.6
Age Risk	1:768		Nuchal translucency MoM	0.80
Biochemical T21 risk	1:2485		Nasal bone	Present
Combined trisomy 21 risk	<1:10000		Sonographer	DR.NIDHI
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 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 hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values</p>	
			<p><b>Trisomy 13/18 + NT</b></p> <p>The calculated risk for Trisomy 13/18 (with NT) is &lt; 1:10000, which indicates a low risk</p>	

 Risk Above Cut Off

 Risk above Age Risk

 Risk below Age risk