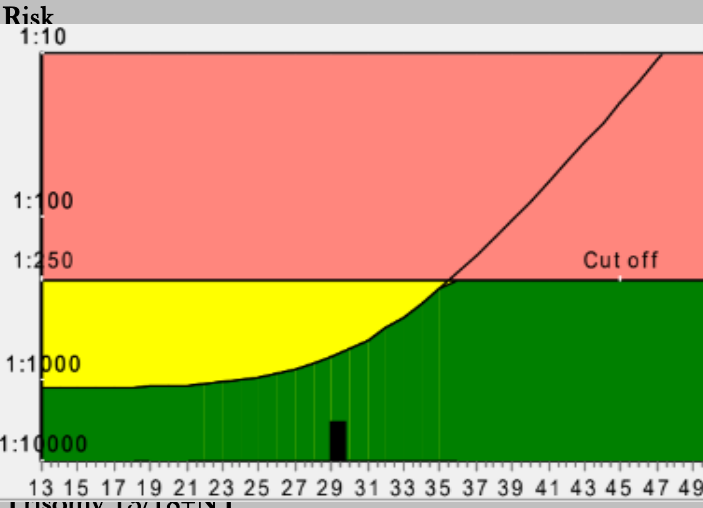


Date of Report 25-02-2024  
PRISCA 5.2.0.13

Patient Data					
Name	MRS. PRAVEEN KUMARI		Patient ID	12402220171	
Birthday	19-10-1994		Sample ID	11829445	
Age at Sample date	54		Sample Date	22-02-2024	
Gestational age	13+0				
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	54	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			
PAPP-A	5.9 mIU/ml	0.80	Gestational age	12+6	
fb-hCG	22.6 ng/ml	0.67	Method	CRL (<>Robinson)	
Risks at sampling date			Scan date	22-02-2024	
Age Risk	1:709		Crown rump length in mm	64.4	
Biochemical T21 risk	1:6522		Nuchal translucency MoM	0.97	
Combined trisomy 21 risk	<1:10000		Nasal bone	absent	
Trisomy 13/18 + NT	<1:10000		Sonographer	DR. AMENDA	
Risk			Qualifications in measuring NT	MD	
			Down's Syndrome Risk (Trisomy 21 Screening)		
<p>The calculated risk for Trisomy 13/18 (with NT) is &lt;1:10000 , which indicates a low risk</p>			<p>The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 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! 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>		

Risk Above Cut Off
  Risk above Age Risk
  Risk below Age risk