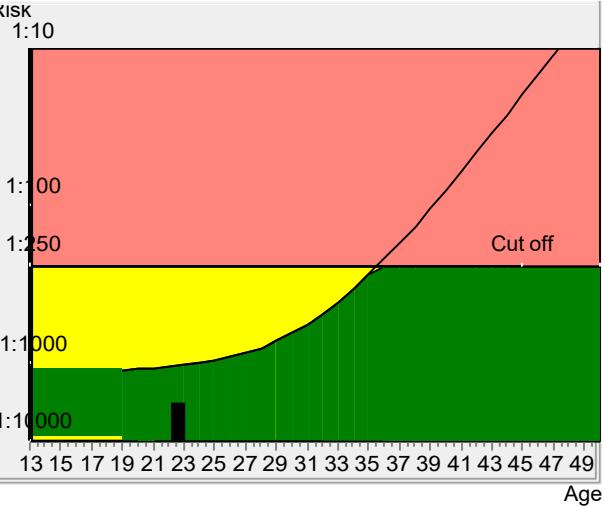


Uma maheshwari devi

Patient data					
Name	Mrs. P.SIRISHA	Patient ID	0012203030318		
Birthday	26-07-1999	Sample ID	23205461		
Age at sample date	22.6	Sample Date	03-03-2022		
Gestational age	13 + 1				
Correction factors					
Fetuses	1	IVF	no		
Weight	38	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data		Ultrasound data			
Parameter	Value	Corr. MoM			
PAPP-A	4.69 mIU/mL	0.55	Gestational age		
fb-hCG	39.42 ng/mL	0.93	Method		
Risks at sampling date		CRL Robinson			
Age risk	1:1062	Scan date			
Biochemical T21 risk	1:1937	Crown rump length in mm			
Combined trisomy 21 risk	1:5513	69.11			
Trisomy 13/18 + NT	<1:10000	Nuchal translucency MoM			
Risk		1.15			
1:10		Nasal bone			
1:100		unknown			
1:250		Sonographer			
1:1000		DR NA			
1:10000		Qualifications in measuring NT			
		MD			
Trisomy 21					
The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.					
After the result of the Trisomy 21 test (with NT) it is expected that among 5513 women with the same data, there is one woman with a trisomy 21 pregnancy and 5512 women with not affected pregnancies.					
The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.					
Please note that risk calculations are statistical approaches and have no diagnostic value!					
The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).					
The laboratory can not be held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!					
					
Trisomy 13/18 + NT					
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.					

Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off