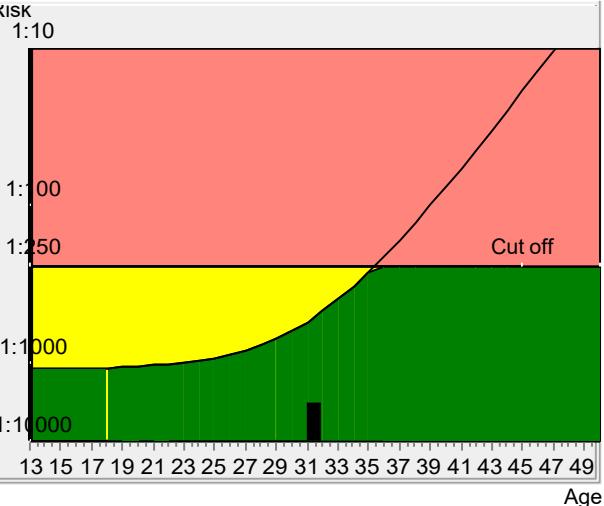


Patient data			
Name	Mrs. K SRUTHI	Patient ID	0012208090135
Birthday	17-03-1991	Sample ID	23226193
Age at sample date	31.4	Sample Date	09-08-2022
Gestational age	12 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	31	diabetes	no
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.1 mIU/mL	0.36	Gestational age 11 + 5
fb-hCG	51.32 ng/mL	0.88	Method CRL Robinson
Risks at sampling date		Scan date 07-08-2022	
Age risk	1:522	Crown rump length in mm 52.8	
Biochemical T21 risk	1:317	Nuchal translucency MoM 0.71	
Combined trisomy 21 risk	1:2033	Nasal bone present	
Trisomy 13/18 + NT	<1:10000	Sonographer NA	
Risk		Qualifications in measuring NT NAGA SAI TEJA	
		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 2033 women with the same data, there is one woman with a trisomy 21 pregnancy and 2032 women with not affected pregnancies. The PAPP-A level is low. 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