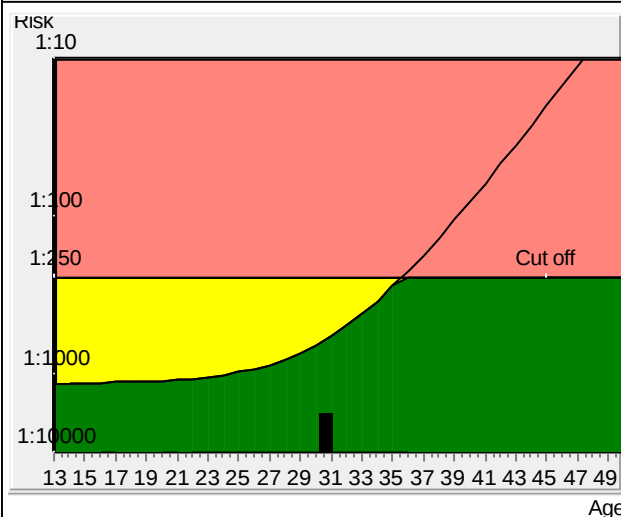


Prisca 5.1.0.17
Date of report: 21-03-2024

Patient data				
Name	Mrs. G POOJITHA SREE	Patient ID	0042403200001	
Birthday	05-08-1993	Sample ID	23124913	
Age at sample date	30.6	Sample Date	20-03-2024	
Gestational age	13 + 4			
Correction factors				
Fetuses	1	IVF	no	
Weight	53	diabetes	no	
Smoker	no	Origin	Asian	
		Previous trisomy 21 pregnancies	unknown	
Biochemical data		Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	13 + 4
PAPP-A	14.15 mIU/mL	2.09	Method	CRL Robinson
fb-hCG	39.54 ng/mL	1.19	Scan date	20-03-2024
Risks at sampling date			Crown rump length in mm	78
Age risk	1:618		Nuchal translucency MoM	0.58
Biochemical T21 risk	1:9399		Nasal bone	present
Combined trisomy 21 risk	<1:10000		Sonographer	NA
Trisomy 13/18 + NT	<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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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 hold 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