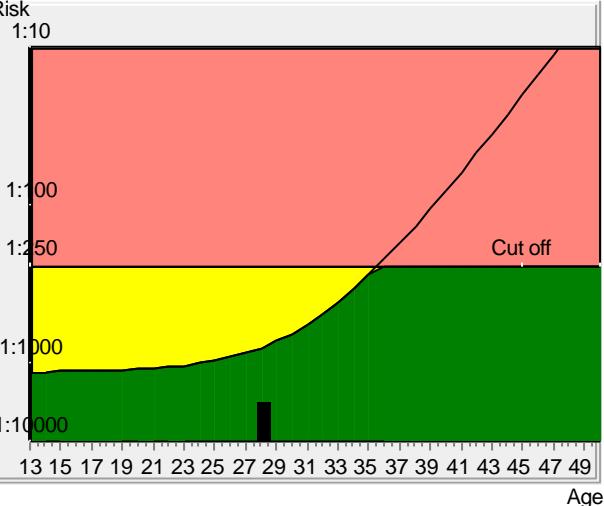


NA

Patient data					
Name	Mrs. POOJITHA	Patient ID	0352411220040		
Birthday	23-08-1996	Sample ID	A1811212		
Age at sample date	28.2	Sample Date	22-11-2024		
Gestational age	13 + 2				
Correction factors					
Fetuses	1	IVF	no		
Weight	67	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data		Ultrasound data			
Parameter	Value	Corr. MoM			
PAPP-A	3.2 mIU/mL	0.59	Gestational age 13 + 2		
fb-hCG	32 ng/ml	1.00	Method CRL Robinson		
Risks at sampling date		Scan date 22-11-2024			
Age risk	1:799	Crown rump length in mm 74			
Biochemical T21 risk	1:1467	Nuchal translucency MoM 0.71			
Combined trisomy 21 risk	1:8361	Nasal bone present			
Trisomy 13/18 + NT	<1:10000	Sonographer NA			
Trisomy 21		Qualifications in measuring NT MD			
		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. <p>After the result of the Trisomy 21 test (with NT) it is expected that among 8361 women with the same data, there is one woman with a trisomy 21 pregnancy and 8360 women with not affected pregnancies.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>			
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