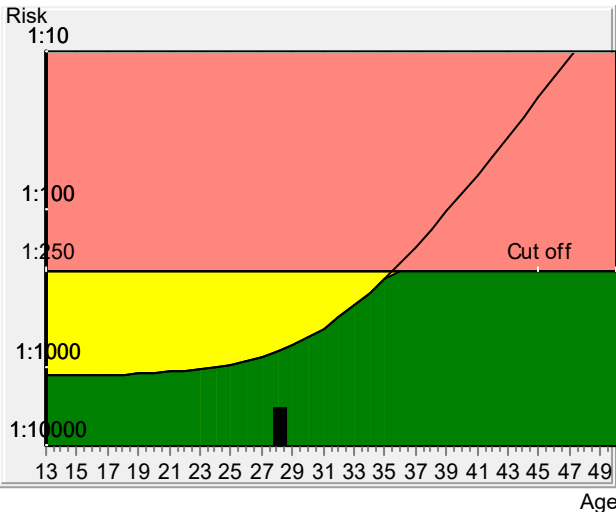


NA

Patient data			
Name		Mrs. TWINKLE BAJAJ	
Patient ID		0622412030066	
Birthday		13-10-1996	
Sample ID		24241663	
Age at sample date		28.1	
Sample Date		03-12-2024	
Gestational age		12 + 6	
Correction factors			
Fetuses	1	IVF	no
Weight	55	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	4.14 mIU/mL	0.73	Method
fb-hCG	21.59 ng/ml	0.57	CRL Robinson
Risks at sampling date			Scan date
			01-12-2024
Age risk			Crown rump length in mm
1:795			63
Biochemical T21 risk			Nuchal translucency MoM
1:7978			0.92
Combined trisomy 21 risk			Nasal bone
<1:10000			present
Trisomy 13/18 + NT			Sonographer
<1:10000			NA
			Qualifications in measuring NT
			MD
Risk			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