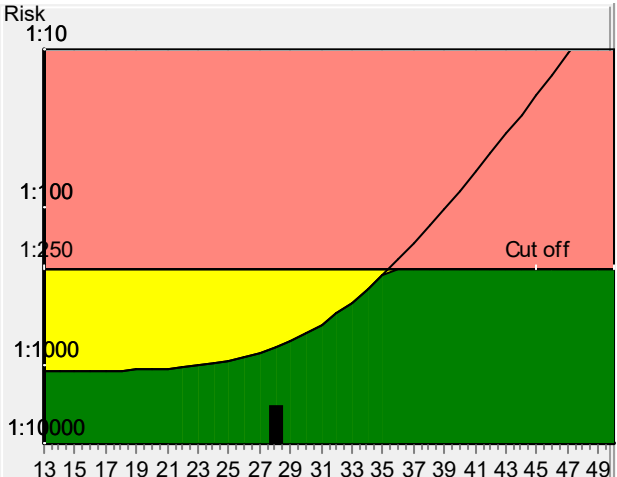


Patient data					
Name		Mrs. SAI SHIRISHA		Patient ID 0012309120285	
Birthday		30/08/95		Sample ID 24332955	
Age at sample date		28.0		Sample Date 12/09/23	
Gestational age		12 + 1			
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	55	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age 12 + 3		
PAPP-A	5.75 mIU/mL	1.54	Method CRL Robinson		
fb-hCG	45.21 ng/mL	0.96	Scan date 14/09/23		
Risks at sampling date			Crown rump length in mm 61.9		
Age risk		1:782	Nuchal translucency MoM 0.81		
Biochemical T21 risk		<1:10000	Nasal bone present		
Combined trisomy 21 risk		<1:10000	Sonographer NA		
Trisomy 13/18 + NT		<1:10000	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

 below cut off
  Below Cut Off, but above Age Risk
  above cut off