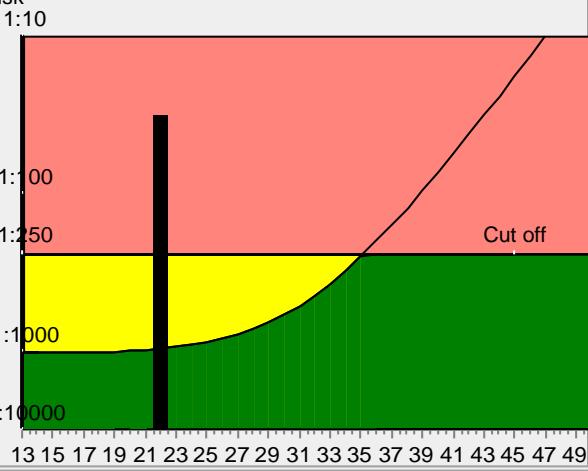


Prisca 5.1.0.17
Date of report: 06-12-2022

Patient data			
Name	Mrs. NIDHI GAUR	Patient ID	0382212030038
Birthday	31-12-2000	Sample ID	24014373
Age at sample date	21.9	Sample Date	03-12-2022
Gestational age	10 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	70	diabetes	no
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.32 mIU/mL	0.95	Gestational age 10 + 5
fb-hCG	55.21 ng/mL	0.96	Method CRL Robinson
Risks at sampling date			
Age risk	1:985		Scan date 03-12-2022
Biochemical T21 risk	1:6014		Crown rump length in mm 40.4
Combined trisomy 21 risk	>1:50		Nuchal translucency MoM 2.62
Trisomy 13/18 + NT	1:326		Nasal bone unknown
Risk		Sonographer NA	
1:10		Qualifications in measuring NT NA	
		Trisomy 21 The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk. After the result of the Trisomy 21 Test (with nuchal translucency), it is expected that among less than 50 pregnancies with the same data, there is one 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 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 1:326, which represents a low risk.	

Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off