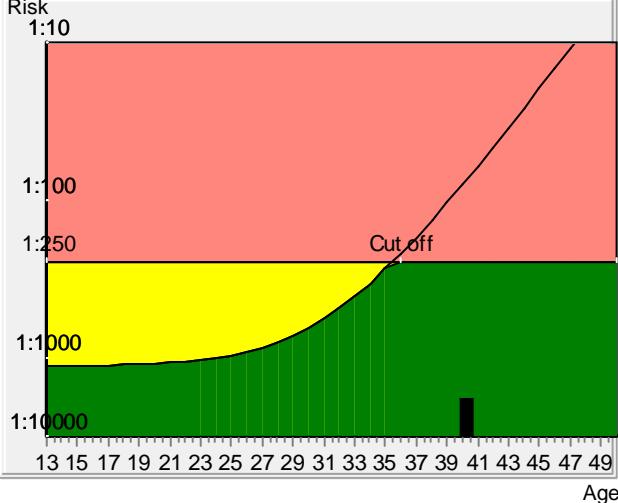


Prisca 5.2.0.13
Date of report: 12/01/22

Patient data				
Name	ASIMA MIDYA	Patient ID		
Birthday	09/10/81	Sample ID		
Age at sample date	40.3	Sample Date		
Gestational age	12 + 5			
Correction factors				
Fetuses	1	IVF		
Weight	71	unknown		
Smoker	unknown	diabetes		
		Origin		
		Asian		
Biochemical data				
Parameter	Value	Corr. MoM		
PAPP-A	3.65 mIU/mL	0.89		
fb-hCG	14.49 ng/mL	0.45		
Risks at sampling date				
Age risk	1:74			
Biochemical T21 risk	1:1934			
Combined trisomy 21 risk	1:8917			
Trisomy 13/18 + NT	<1:10000			
Ultrasound data				
Gestational age	12 + 4			
Method	CRL	Robinson		
Scan date	09/01/22			
Crown rump length in mm	63.1			
Nuchal translucency MoM	0.84			
Nasal bone	unknown			
Sonographer	DR.ANINDYA BANDYOPADHYAY			
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 8917 women with the same data, there is one woman with a trisomy 21 pregnancy and 8916 women with not affected pregnancies.				
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 < 1:10000, which represents a low risk.				
				

Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off