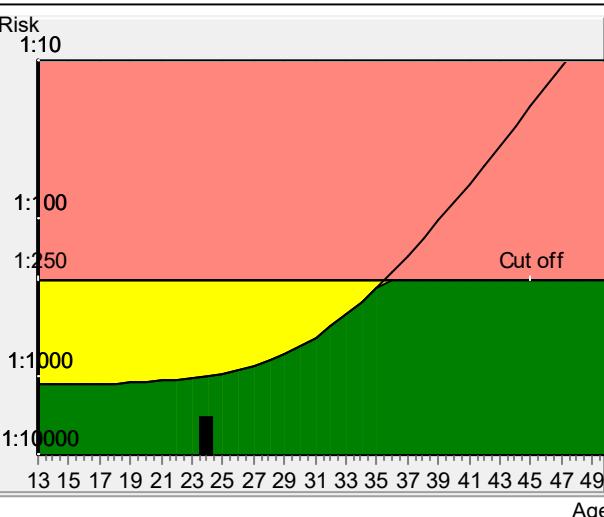


Patient data								
Name	Mrs. SHEETAL DHENGDE	Patient ID	0372205110054					
Birthday	17-06-1998	Sample ID	23369777					
Age at sample date	23.9	Sample Date	11-05-2022					
Gestational age	12 + 6							
Correction factors								
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	40	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	6.45 mIU/mL	0.90	Gestational age 12 + 5					
fb-hCG	48.66 ng/mL	1.08	Method CRL Robinson					
Risks at sampling date								
Age risk	1:1014		Scan date 10-05-2022					
Biochemical T21 risk	1:4155		Crown rump length in mm 65.9					
Combined trisomy 21 risk	1:5060		Nuchal translucency MoM 1.37					
Trisomy 13/18 + NT	<1:10000		Nasal bone unknown					
Sonographer DR NA								
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 5060 women with the same data, there is one woman with a trisomy 21 pregnancy and 5059 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