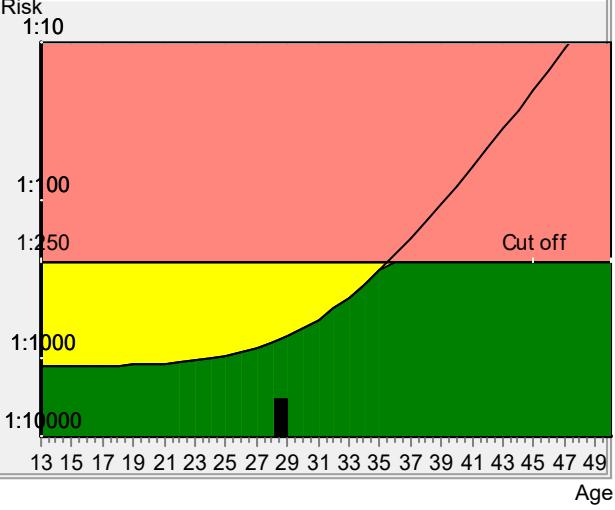


Prisca 5.1.0.17

Date of report: 30/01/26

Patient data								
Name	Mrs. VAISHALI NAGE	Patient ID	0672601290248					
Birthday	10/06/97	Sample ID	Mrs. VAISHALI NAGE					
Age at sample date	28.6	Sample Date	29/01/26					
Gestational age	13 + 0							
Correction factors								
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	47	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	3.66 mIU/mL	0.58	Gestational age 12 + 6					
fb-hCG	36.95 ng/mL	0.91	Method CRL Robinson					
Risks at sampling date								
Age risk	1:763		Scan date 28/01/26					
Biochemical T21 risk	1:1609		Crown rump length in mm 67.2					
Combined trisomy 21 risk	1:6942		Nuchal translucency MoM 1.00					
Trisomy 13/18 + NT	<1:10000		Nasal bone present					
Sonographer NA								
Qualifications in measuring NT Sonographer								
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 6942 women with the same data, there is one woman with a trisomy 21 pregnancy and 6941 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