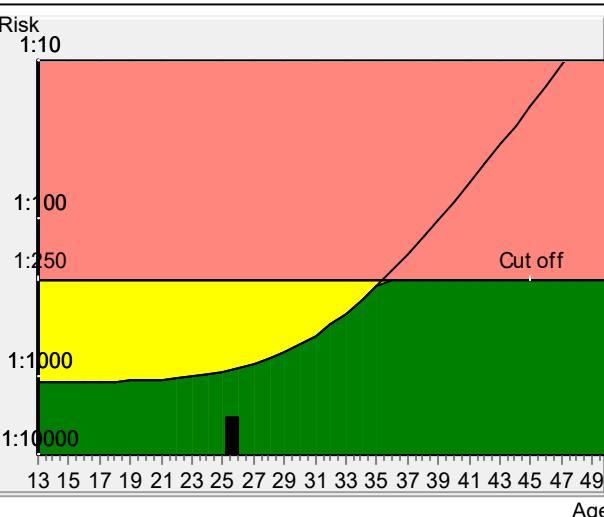


Prisca 5.1.0.17

Date of report: 23/10/25

NA

Patient data								
Name	Ms. JHARANA SWAIN	Patient ID	0642510190068					
Birthday	10/03/00	Sample ID	B3897841					
Age at sample date	25.6	Sample Date	19/10/25					
Gestational age	12 + 1							
Correction factors								
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	48	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	2.98 mIU/mL	0.68	Gestational age 12 + 1					
fb-hCG	42.46 ng/mL	0.86	Method CRL Robinson					
Risks at sampling date								
Age risk	1:921		Scan date 19/10/25					
Biochemical T21 risk	1:3315		Crown rump length in mm 58.6					
Combined trisomy 21 risk	<1:10000		Nuchal translucency MoM 0.78					
Trisomy 13/18 + NT	<1:10000		Nasal bone present					
Sonographer NA								
Qualifications in measuring NT NA								
Trisomy 21								
The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. <p>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.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>								
								
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