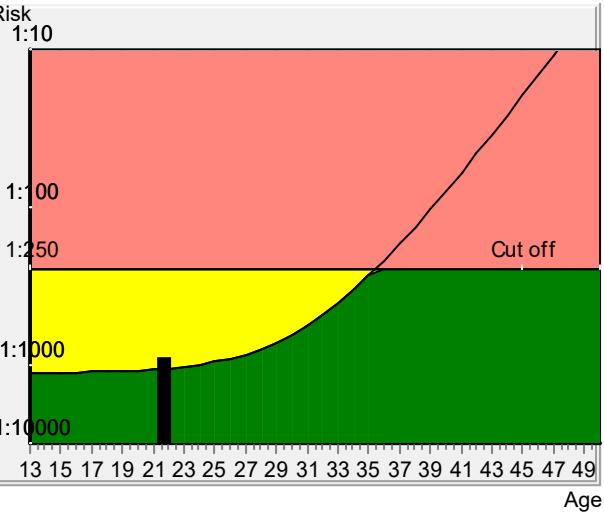


Prisca 5.1.0.17

Date of report: 22/11/25

Patient data									
Name		Patient ID		0312511210040					
Birthday		Sample ID		A0794219					
Age at sample date		Sample Date		21/11/25					
Gestational age		12 + 4							
Correction factors									
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown				
Weight	67	diabetes	no						
Smoker	no	Origin	Asian						
Biochemical data									
Parameter	Value	Corr. MoM	Ultrasound data						
PAPP-A	1.22 mIU/mL	0.34	Gestational age 12 + 4						
fb-hCG	41.7 ng/mL	1.04	Method CRL Robinson						
Risks at sampling date									
Age risk		1:1062	Scan date 21/11/25						
Biochemical T21 risk		1:398	Crown rump length in mm 63						
Combined trisomy 21 risk		1:896	Nuchal translucency MoM 1.23						
Trisomy 13/18 + NT		1:9108	Nasal bone present						
Risk 1:10		Sonographer NA		Sonographer					
		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. <p>After the result of the Trisomy 21 test (with NT) it is expected that among 896 women with the same data, there is one woman with a trisomy 21 pregnancy and 895 women with not affected pregnancies.</p> <p>The PAPP-A level is low.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.</p> <p>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 1:9108, which represents a low risk.									

Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off