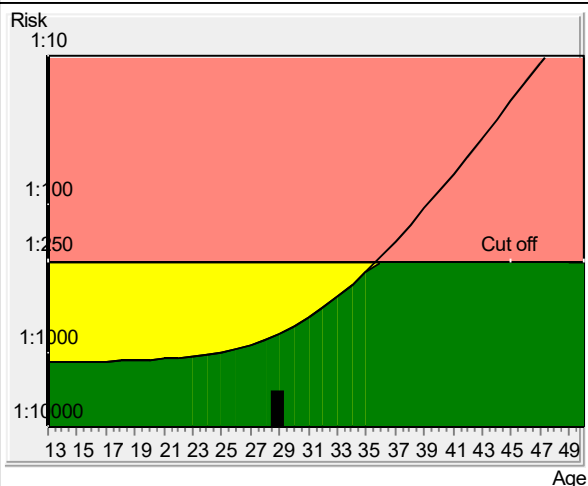


Prisca 5.2.0.13

Date of report: 16/04/25

Patient data			
Name	MRS. NANCY		Patient ID
Birthday	19/05/96		Sample ID
Age at sample date	28.9		Sample Date
Gestational age	13 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	48.7	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		no	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	10.1 mIU/ml	1.24	13 + 5
fb-hCG	46.4 ng/ml	1.10	Method
Risks at sampling date			CRL Robinson
Age risk	1:761		Scan date
Biochemical T21 risk	1:5969		14/04/25
Combined trisomy 21 risk	<1:10000		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		80.5
			Nuchal translucency MoM
			0.67
			Nasal bone
			present
			Sonographer
			..
			Qualifications in measuring NT
			..
Trisomy 21			
<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <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 hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>			
Trisomy 13/18 + NT			
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			



Sign of Physician

below cut off
 Below Cut Off, but above Age Risk
 above cut off