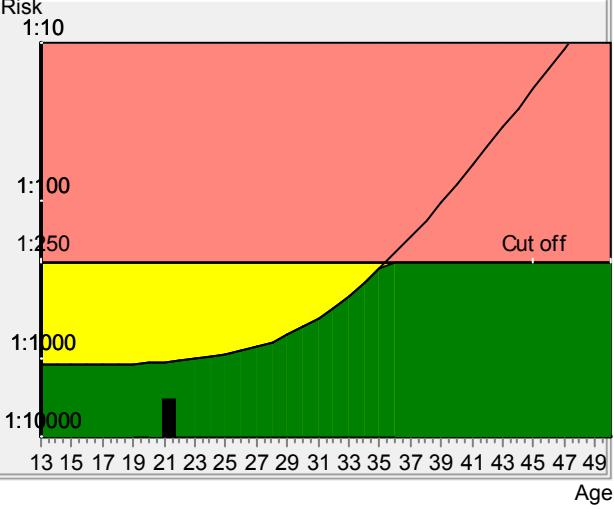


Prisca 5.2.0.13

Date of report: 05/10/25

Dr

XYZ

Patient data			
Name	AMRITA	Patient ID	2M036598
Birthday	02/06/04	Sample ID	2M036598
Age at sample date	21.3	Sample Date	04/10/25
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	unknown
Weight	50	diabetes	unknown
Smoker	unknown	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	6.39 mIU/mL	1.22	Gestational age 12 + 2
fb-hCG	35 ng/mL	0.86	Method CRL Robinson
Risks at sampling date			
Age risk	1:1059		Scan date 04/10/25
Biochemical T21 risk	<1:10000		Crown rump length in mm 59
Combined trisomy 21 risk	<1:10000		Nuchal translucency MoM 0.78
Trisomy 13/18 + NT	<1:10000		Nasal bone unknown
Sonographer DR SELF			
Qualifications in measuring NT MD			
Trisomy 21			
<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></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 held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>			
			
Trisomy 13/18 + NT			
<p><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>			

Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off