

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
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	6.39 mIU/mL	1.22	12 + 2
fb-hCG	35 ng/mL	0.86	Method
			CRL Robinson
			Scan date
			04/10/25
Risks at sampling date		Crown rump length in mm	
Age risk	1:1059	59	
Biochemical T21 risk	<1:10000	Nuchal translucency MoM	
Combined trisomy 21 risk	<1:10000	0.78	
Trisomy 13/18 + NT	<1:10000	Nasal bone	
		unknown	
		Sonographer	
		DR SELF	
		Qualifications in measuring NT	
		MD	
Risk		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! 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 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