

Prisca 5.1.0.17
Date of report: 25/01/26

Patient data			
Name	Mrs. AYESHA	Patient ID	0352601240048
Birthday	17/01/89	Sample ID	b3452972
Age at sample date	37.0	Sample Date	23/01/26
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	58	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.86 mIU/mL	0.52	Gestational age 13 + 1
fb-hCG	34.02 ng/mL	0.98	Method CRL Robinson
			Scan date 22/01/26
Risks at sampling date			Trisomy 21
Age risk		1:175	Crown rump length in mm 71
Biochemical T21 risk		1:239	Nuchal translucency MoM 0.79
Combined trisomy 21 risk		1:1409	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer NA
			Qualifications in measuring NT Sonographer
			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 1409 women with the same data, there is one woman with a trisomy 21 pregnancy and 1408 women with not affected pregnancies. 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!
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