

Prisca 5.2.0.13
Date of report: 31/08/2025

Patient data			
Name	MRS. FARNAZ KHATOON		Patient ID
Birthday	17/05/1993		Sample ID
Age at sample date	32.3		Sample Date
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	58	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		no	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.26 mIU/ml	0.64	11 + 0
fb-hCG	40.7 ng/ml	0.95	Method
Risks at sampling date			CRL Robinson
Age risk	1:470		Scan date
Biochemical T21 risk	1:1185		16/08/2025
Combined trisomy 21 risk	1:642		Crown rump length in mm
Trisomy 13/18 + NT	1:9693		43.3
			Nuchal translucency MoM
			1.58
			Nasal bone
			present
			Sonographer
			DR. SYED ZAIN ABBAS
			Qualifications in measuring NT
			M.D
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 642 women with the same data, there is one woman with a trisomy 21 pregnancy and 641 women with not affected pregnancies.</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 1:9693, which represents a low risk.</p>			

Sign of Physician