

Prisca 5.1.0.17
Date of report: 27-01-2026

Patient data			
Name	Mrs. MARVA SHARIQ	Patient ID	0692601220110
Birthday	21-08-1995	Sample ID	D0038512
Age at sample date	30.4	Sample Date	21-01-2026
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	65	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	0.96 mIU/mL	0.25	12 + 0
fb-hCG	40.52 ng/mL	1.04	Method
			CRL Robinson
			Scan date
			16-01-2026
Risks at sampling date			Trisomy 21
Age risk		1:616	The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
Biochemical T21 risk		1:84	After the result of the Trisomy 21 test (with NT) it is expected that among 320 women with the same data, there is one woman with a trisomy 21 pregnancy and 319 women with not affected pregnancies.
Combined trisomy 21 risk		1:320	The PAPP-A level is low.
Trisomy 13/18 + NT		1:1920	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 1:1920, which represents a low risk.			

Sign of Physician