

Prisca 5.1.0.17
Date of report: 01-12-2025

Patient data			
Name	Mrs. SWETA SINGH	Patient ID	0622511260047
Birthday	05-10-1996	Sample ID	B2801238
Age at sample date	29.1	Sample Date	26-11-2025
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	63	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.76 mIU/mL	0.39	12 + 6
fb-hCG	38.69 ng/mL	1.06	Method
			CRL Robinson
			Scan date
			25-11-2025
Risks at sampling date		Trisomy 21	
Age risk	1:725	Crown rump length in mm	68
Biochemical T21 risk	1:382	Nuchal translucency MoM	0.76
Combined trisomy 21 risk	1:2452	Nasal bone	unknown
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 2452 women with the same data, there is one woman with a trisomy 21 pregnancy and 2451 women with not affected pregnancies. The PAPP-A level is low. 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