

Prisca 5.1.0.17
Date of report: 10-06-2025

NA

Patient data			
Name	Mrs. ARCHANA KUMARI	Patient ID	0482506080217
Birthday	23-06-1999	Sample ID	B2538246
Age at sample date	26.0	Sample Date	08-06-2025
Gestational age	11 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	53	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.76 mIU/mL	0.86	11 + 4
fb-hCG	47.08 ng/mL	0.90	Method
			CRL Robinson
			Scan date
			07-06-2025
Risks at sampling date			
Age risk		1:890	Crown rump length in mm
Biochemical T21 risk		1:4946	50.93
Combined trisomy 21 risk		1:1723	Nuchal translucency MoM
Trisomy 13/18 + NT		<1:10000	1.68
			Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			NA
Trisomy 21			
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 1723 women with the same data, there is one woman with a trisomy 21 pregnancy and 1722 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
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