

Prisca 5.1.0.17
Date of report: 15/11/25

NA

Patient data				
Name	Mrs. MANISHA SINGH		Patient ID	0502511130005
Birthday	24/12/92		Sample ID	B3852353
Age at sample date	32.9		Sample Date	13/11/25
Gestational age	13 + 3			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	58	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 0
PAPP-A	1.13 mIU/mL	0.20	Method	CRL Robinson
fb-hCG	32.8 ng/mL	0.98	Scan date	03/11/25
Risks at sampling date			Crown rump length in mm	
Age risk	1:429		55.77	
Biochemical T21 risk	>1:50		Nuchal translucency MoM	
Combined trisomy 21 risk	>1:50		1.44	
Trisomy 13/18 + NT	1:88		Nasal bone	
			present	
			Sonographer	
			NA	
			Qualifications in measuring NT	
			NA	
Trisomy 21				
The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk. After the result of the Trisomy 21 Test (with nuchal translucency), it is expected that among less than 50 pregnancies with the same data, there is one trisomy 21 pregnancy. 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 test (with nuchal translucency) is 1:88, which represents an increased risk.				

Sign of Physician

below cut off	Below Cut Off, but above Age Risk	above cut off
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