

Date of report: 20-07-2024

N A

Patient data				
Name	Mrs. S.SRIVANI		Patient ID	0352407190020
Birthday	27-06-2001		Sample ID	A0228867
Age at sample date	23.1		Sample Date	19-07-2024
Gestational age	12 + 5			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21
Weight	37	diabetes	no	pregnancies
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 5
PAPP-A	10.25 mIU/mL	1.38	Method	CRL Robinson
fb-hCG	48.96 ng/mL	1.02	Scan date	19-07-2024
Risks at sampling date			Crown rump length in mm	66
Age risk	1:1035		Nuchal translucency MoM	0.77
Biochemical T21 risk	<1:10000		Nasal bone	present
Combined trisomy 21 risk	<1:10000		Sonographer	N A
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	MD
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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