

Date of report: 26-08-2024

N A

Patient data				
Name	Mrs. MINAKSHI RAI		Patient ID	0482408230329
Birthday	23-08-1995		Sample ID	A0637956
Age at sample date	29.0		Sample Date	23-08-2024
Gestational age	13 + 1			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21
Weight	63	diabetes	no	pregnancies
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	13 + 0
PAPP-A	1.77 mIU/mL	0.37	Method	CRL Robinson
fb-hCG	37.41 ng/mL	1.06	Scan date	22-08-2024
Risks at sampling date			Crown rump length in mm	70
Age risk		1:739	Nuchal translucency MoM	0.74
Biochemical T21 risk		1:337	Nasal bone	present
Combined trisomy 21 risk		1:2194	Sonographer	N A
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD
RISK			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 2194 women with the same data, there is one woman with a trisomy 21 pregnancy and 2193 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

below cut off



Below Cut Off, but above Age Risk



above cut off