

N A

Patient data				
Name	Mrs. POOJA NAYAK		Patient ID	0662408160064
Birthday	11-04-1992		Sample ID	A0970035
Age at sample date	32.3		Sample Date	16-08-2024
Gestational age	13 + 3			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21
Weight	102	diabetes	no	pregnancies
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 4
PAPP-A	3.37 mIU/mL	1.18	Method	CRL Robinson
fb-hCG	43.69 ng/mL	1.53	Scan date	10-08-2024
Risks at sampling date			Crown rump length in mm	64
Age risk	1:472		Nuchal translucency MoM	1.22
Biochemical T21 risk	1:1538		Nasal bone	present
Combined trisomy 21 risk	1:3117		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 3117 women with the same data, there is one woman with a trisomy 21 pregnancy and 3116 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