

Patient data			
Name	MRS. BHARTI	Patient ID	
Birth day	15-10-1996	Sample ID	2305028688/NOD
Age at sample date	26.6	Sample Date	15-05-2023
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	60	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		no	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.88 mIU/ml	1.63	Gestational age 12 + 0
fb-hCG	144 ng/ml	3.12	Method CRL Robinson
Risks at sampling date			Scan date 13-05-2023
Age risk		1:876	Crown rump length in mm 56.7
Biochemical T21 risk		1:825	Nuchal translucency MoM 0.54
Combined trisomy 21 risk		1:4241	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer ..
Risk			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 4241 women with the same data, there is one woman with a trisomy 21 pregnancy and 4240 women with not affected pregnancies.
			The free beta HCG level is high.
			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.



