

Patient data			
Name	MRS. POONAM	Patient ID	
Birthday	01-01-1995	Sample ID	2305050734/NOD
Age at sample date	28.4	Sample Date	25-05-2023
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	50	diabetes	no
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	5.95 mIU/ml	0.84	13 + 2
fb-hCG	132.4 ng/ml	3.07	Method
Risks at sampling date			CRL Robinson
Age risk Biochemical	1:792		Scan date
T21 risk	1:209		24-05-2023
Combined trisomy 21 risk	1:1272		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		74
Risk	1:10		Nuchal translucency MoM
	1:100		0.60
	1:250		Nasal bone
	1:1000		present
	1:10000		Sonographer
	1:100000		..
	13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49		Qualifications in measuring NT
	Age		MD

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 1272 women with the same data, there is one woman with a trisomy 21 pregnancy and 1271 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 held 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.



