

Patient data			
Name	MRS. KAVITA	Patient ID	
Birthday	03-08-2000	Sample ID	2305062230/NOD
Age at sample date	22.8	Sample Date	31-05-2023
Gestational age	12 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	58	diabetes	no
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.66 mIU/ml	0.81	11 + 6
fb-hCG	81.6 ng/ml	1.69	Method
Risks at sampling date			CRL Robinson
Age risk	1:1016		Scan date
Biochemical T21 risk	1:1152		30-05-2023
Combined trisomy 21 risk	1:5915		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		55
Risk	1:10		Nuchal translucency MoM
	1:100		0.89
	1:250		Nasal bone
			present
	1:1000		Sonographer
	1:10000		..
	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
			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 5915 women with the same data, there is one woman with a trisomy 21 pregnancy and 5914 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.