

Patient data		MRS. MANA BI	
Name	22-05-1998	Patient ID	
Birth day		Sample ID	2306011700/NOD
Age at sample date	25.0	Sample Date	06-06-2023
Gestational age	12 + 6		
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	5.16 mIU/ml	0.91	12 + 5
fb-hCG	44.2 ng/ml	0.96	Method
Risks at sampling date			CRL Robinson
Age risk	1:971		Scan date
Biochemical T21 risk	1:5339		05-06-2023
Combined trisomy 21 risk	<1:10000		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		66
Risk			Nuchal translucency MoM
			0.59
			Nasal bone
			present
			Sonographer
			..
			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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.
			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.

