

NA

Patient data			
Name	Mrs. MANJULA	Patient ID	0352412100004
Birthday	20/06/03	Sample ID	a1251753
Age at sample date	21.5	Sample Date	10/12/24
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	41	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	9.61 mIU/mL	1.12	13 + 2
fb-hCG	32.64 ng/mL	0.86	Method
			CRL Robinson
			Scan date
			09/12/24
Risks at sampling date			Crown rump length in mm
Age risk		1:1097	73.1
Biochemical T21 risk		<1:10000	Nuchal translucency MoM
Combined trisomy 21 risk		<1:10000	0.99
Trisomy 13/18 + NT		<1:10000	Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			MD
Risk			Trisomy 21
1:10			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
1:100			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.
1:250			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!
1:1000			The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).
1:10000			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