

Patient data			
Name	Mrs. KUMARI N.M	Patient ID	0662312210053
Birthday	31-10-1996	Sample ID	25106235
Age at sample date	27.1	Sample Date	21-12-2023
Gestational age	12 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	61	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	8.65 mIU/mL	2.78	11 + 6
fb-hCG	45.21 ng/mL	0.97	Method
			CRL Robinson
			Scan date
			20-12-2023
Risks at sampling date			Crown rump length in mm
Age risk		1:835	54.5
Biochemical T21 risk		<1:10000	Nuchal translucency MoM
Combined trisomy 21 risk		<1:10000	0.97
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 PAPP-A level is high.		
1:1000	The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.		
1:10000	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!		
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49	Age		
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