

Patient data			
Name	Mrs. NIDHI SHRIWASTAV	Patient ID	0482402250173
Birthday	26-03-1985	Sample ID	A0059663
Age at sample date	38.9	Sample Date	25-02-2024
Gestational age	13 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	74	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	4.98 mIU/mL	1.05	12 + 5
fb-hCG	29.22 ng/mL	1.02	Method
Risks at sampling date			CRL Robinson
Age risk	1:110		Scan date
Biochemical T21 risk	1:712		18-02-2024
Combined trisomy 21 risk	1:3518		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		65.7
			Nuchal translucency MoM
			0.36
			Nasal bone
			present
			Sonographer
			N A
			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 3518 women with the same data, there is one woman with a trisomy 21 pregnancy and 3517 women with not affected pregnancies.	
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!	
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