


Patient data			
Name	MRS. SURBHI KUMARI	Patient ID	
Birthday	15/08/2001	Sample ID	2312003655/NOD
Age at sample date	22.3	Sample Date	3/12/2023
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	67	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.7 mIU/ml	0.71	12 + 4
fb-hCG	37.1 ng/ml	0.88	Method
			CRL Robinson
Risks at sampling date			Scan date
Age risk	1:1055		2/12/2023
Biochemical T21 risk	1:4040		Crown rump length in mm
Combined trisomy 21 risk	<1:10000		63.2
Trisomy 13/18 + NT	<1:10000		Nuchal translucency MoM
			0.68
			Nasal bone
			present
			Sonographer
			DR. JAIDEEP TOMAR
			Qualifications in measuring NT
			M.D
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!	
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

 below cut off	 Below Cut Off, but above Age Risk	 above cut off
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