

Patient data			
Name	MRS. SMARTHA	Patient ID	
Birthday	02-05-1991	Sample ID	2111220340/AMB
Age at sample date	30.5	Sample Date	12-11-2021
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	yes
Weight	66.2	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	0.74	12 + 1
fb-hCG	77.1 ng/ml	1.78	Method
			CRL Robinson
Risks at sampling date			Scan date
Age risk		1:604	09-11-2021
Biochemical T21 risk		1:495	Crown rump length in mm
Combined trisomy 21 risk		1:2121	57.7
Trisomy 13/18 + NT		<1:10000	Nuchal translucency MoM
			0.99
			Nasal bone
			present
			Sonographer
			DR. SANJEEV SHARMA
			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 2121 women with the same data, there is one woman with a trisomy 21 pregnancy and 2120 women with not affected pregnancies.	
1:250		The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.	
1:1000		Please note that risk calculations are statistical approaches and have no diagnostic value!	
1:10000		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



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