

Patient data			
Name	MRS, SHREYA BHAT	Patient ID	4115664
Birth day	07-09-1991	Sample ID	4115664
Age at delivery	32.0	Sample Date	16-02-2023
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	unknown
Weight	70	diabetes	unknown
Smoker	unknown	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.9 mIU/ml	0.65	12 + 2
fb-hCG	23.5 ng/ml	0.70	Method
			CRL Robinson
			Scan date
			14-02-2023
Risks at term			Crown rump length in mm
Age risk		1:743	60.4
Biochemical T21 risk		1:3623	Nuchal translucency MoM
Combined trisomy 21 risk		<1:10000	1.15
Trisomy 13/18 + NT		<1:10000	Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			FMF
Risk	Trisomy 21		
1:10	<b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b>		
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.		
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		
	<b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b>		

**Sign of Physician**

