

Patient data			
Name	0012202210167	Patient ID	Mrs. SHAILAJA
Birthday	06/03/03	Sample ID	23049091
Age at sample date	19.0	Sample Date	21/02/22
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	52	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.11 mIU/mL	0.66	
fb-hCG	48.65 ng/mL	1.35	
Risks at sampling date		Gestational age	12 + 6
Age risk	1:1128	Method	CRL Robinson
Biochemical T21 risk	1:1356	Scan date	18/02/22
Combined trisomy 21 risk	1:5771	Crown rump length in mm	67
Trisomy 13/18 + NT	<1:10000	Nuchal translucency MoM	1.00
		Nasal bone	present
		Sonographer	DR AMER NAVEED
		Qualifications in measuring NT	MD
Risk		Trisomy 21	
		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 5771 women with the same data, there is one woman with a trisomy 21 pregnancy and 5770 women with not affected pregnancies. 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! 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!	
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