

Patient data				
Name	Mrs. KHAIRUNNISA NISHA		Patient ID	0012208020190
Birthday	22/03/03		Sample ID	23567392
Age at sample date	19.4		Sample Date	02/08/22
Gestational age	12 + 4			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	52.2	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 3
PAPP-A	2 mIU/mL	0.42	Method	CRL Robinson
fb-hCG	45.22 ng/mL	1.04	Scan date	01/08/22
Risks at sampling date			Crown rump length in mm	
Age risk	1:1098		62.22	
Biochemical T21 risk	1:734		Nuchal translucency MoM	
Combined trisomy 21 risk	1:4613		0.50	
Trisomy 13/18 + NT	<1:10000		Nasal bone	
			unknown	
			Sonographer	
			NA	
			Qualifications in measuring NT	
			NAGA SAI TEJA	
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 4613 women with the same data, there is one woman with a trisomy 21 pregnancy and 4612 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