

Patient data			
Name	MRS. BALJEET KAUR	Patient ID	2108220153/AMB
Birthday	05-02-1994	Sample ID	2108220153/AMB
Age at sample date	27.5	Sample Date	04-08-2021
Gestational age	11 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	55	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.26 mIU/ml	1.16	11 + 3
fb-hCG	155 ng/ml	3.01	Method
			CRL Robinson
			Scan date
			03-08-2021
Risks at sampling date			Crown rump length in mm
Age risk		1:800	49.6
Biochemical T21 risk		1:440	Nuchal translucency MoM
Combined trisomy 21 risk		1:2454	0.68
Trisomy 13/18 + NT		<1:10000	Nasal bone
			present
			Sonographer
			.
			Qualifications in measuring NT
			MD
Risk		Trisomy 21	
		<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 2454 women with the same data, there is one woman with a trisomy 21 pregnancy and 2453 women with not affected pregnancies. The free beta HCG level is high.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>	
Trisomy 13/18 + NT			
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician