

Patient data			
Name	MRS. MANDEEP	Patient ID	2107220888/AMB
Birth day	13-05-1994	Sample ID	2107220888/AMB
Age at sample date	27.2	Sample Date	28-07-2021
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	52.6	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	11.2 mIU/ml	2.09	12 + 5
fb-hCG	108 ng/ml	2.39	Method
			CRL Robinson
Risks at sampling date			Scan date
Age risk		1:857	27-07-2021
Biochemical T21 risk		1:2307	Crown rump length in mm
Combined trisomy 21 risk		<1:10000	65.7
Trisomy 13/18 + NT		<1:10000	Nuchal translucency MoM
			0.45
			Nasal bone
			present
			Sonographer
			.
			Qualifications in measuring NT
			MD
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 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			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician

