

Patient data			
Name	MRS. POOJA	Patient ID	
Birthday	22/01/1996	Sample ID	2406220858/AMB
Age at sample date	28.4	Sample Date	18/06/2024
Gestational age	11 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	38	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.9 mIU/ml	0.79	10 + 6
fb-hCG	78.5 ng/ml	1.28	Method
			CRL Robinson
			Scan date
			15/06/2024
Risks at sampling date		Crown rump length in mm	
Age risk	1:731	42.2	
Biochemical T21 risk	1:1529	Nuchal translucency MoM	
Combined trisomy 21 risk	1:8379	0.56	
Trisomy 13/18 + NT	<1:10000	Nasal bone	
		present	
		Sonographer	
		..	
		Qualifications in measuring NT	
		..	
Risk		Trisomy 21	
1:10		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 8379 women with the same data, there is one woman with a trisomy 21 pregnancy and 8378 women with not affected pregnancies.	
1:100		The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.	
1:250		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)).	
1:1000		The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
1:10000			
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

