

Patient data			
Name	MRS. MANPREET	Patient ID	
Birthday	04-09-1994	Sample ID	2109220735/AMB
Age at sample date	27.1	Sample Date	23-09-2021
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	64	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.8 mIU/ml	0.84	Gestational age 11 + 4
fb-hCG	84.1 ng/ml	1.86	Method CRL Robinson
Risks at sampling date			Scan date 18-09-2021
Age risk		1:849	Crown rump length in mm 51.3
Biochemical T21 risk		1:825	Nuchal translucency MoM 0.94
Combined trisomy 21 risk		1:3858	Nasal bone present
Trisomy 13/18 + NT		<1:10000	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 3858 women with the same data, there is one woman with a trisomy 21 pregnancy and 3857 women with not affected pregnancies.</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