

Patient data			
Name	MRS. MANINDER	Patient ID	1903220275/AMB
Birthday	28-06-1985	Sample ID	1903220275/AMB
Age at sample date	33.7	Sample Date	08-03-2019
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	93	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.5 mIU/ml	1.54	12 + 6
fb-hCG	42.1 ng/ml	1.43	Method
			CRL Robinson
			Scan date
			08-03-2019
Risks at sampling date		Crown rump length in mm	
Age risk	1:361	68	
Biochemical T21 risk	1:2280	Nuchal translucency MoM	
Combined trisomy 21 risk	<1:10000	0.70	
Trisomy 13/18 + NT	<1:10000	Nasal bone	
		present	
		Sonographer	
		Qualifications in measuring NT	
		MD	
Risk		Trisomy 21	
1:10		<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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</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