

Patient data			
Name	Ms.KIRANJEET	Patient ID	211903050007
Birthday	23/12/1987	Sample ID	211903050007
Age at sample date	31.0	Sample Date	05-03-2019
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	unknown
Weight	82	diabetes	unknown
Smoker	unknown	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	4.5 mIU/ml	1.38	12 + 3
fb-hCG	105.1 ng/ml	3.03	Method
			CRL Robinson
			Scan date
			04-03-2018
Risks at sampling date			Crown rump length in mm
Age risk		1:500	59.0
Biochemical T21 risk		1:380	Nuchal translucency MoM
Combined trisomy 21 risk		1:1990	0.78
Trisomy 13/18 + NT		<1:10000	Nasal bone
			present
			Sonographer
			.
			Qualifications in measuring NT
			MD
Risk			Trisomy 21
1:10			<b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b>
			After the result of the Trisomy 21 test (with NT) it is expected that among 1990 women with the same data, there is one woman with a trisomy 21 pregnancy and 1989 women with not affected pregnancies. The free beta HCG level is high. 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! 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!
Trisomy 13/18 + NT			
<b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b>			

Sign of Physician