

Patient data			
Name	MRS. SIMRANJIT	Patient ID	1906220412/AMB
Birthday	21-06-1991	Sample ID	1906220412/AMB
Age at sample date	28.0	Sample Date	11-06-2019
Gestational age	11 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	42	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	8.21 mIU/ml	2.69	11 + 1
fb-hCG	191 ng/ml	3.47	Method
			CRL Robinson
Risks at sampling date		Scan date	10-06-2019
Age risk	1:760	Crown rump length in mm	46
Biochemical T21 risk	1:748	Nuchal translucency MoM	0.95
Combined trisomy 21 risk	1:3035	Nasal bone	present
Trisomy 13/18 + NT	<1:10000	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 3035 women with the same data, there is one woman with a trisomy 21 pregnancy and 3034 women with not affected pregnancies. The free beta HCG level is high. The PAPP-A 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!</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