

Patient data			
Name	MRS. SARBJEET KAUR	Patient ID	1906221129/AMB
Birthday	23-08-1990	Sample ID	1906221129/AMB
Age at sample date	28.8	Sample Date	24-06-2019
Gestational age	11 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	61	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.5 mIU/ml	0.62	11 + 1
fb-hCG	150 ng/ml	3.30	Method
			CRL Robinson
			Scan date
			20-06-2019
Risks at sampling date		Crown rump length in mm	
Age risk	1:714	46.2	
Biochemical T21 risk	1:76	Nuchal translucency MoM	
Combined trisomy 21 risk	1:504	0.79	
Trisomy 13/18 + NT	<1:10000	Nasal bone	
		present	
		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 504 women with the same data, there is one woman with a trisomy 21 pregnancy and 503 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!</p>	
		<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</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

