

Patient data			
Name	MRS. GURWINDER KAUR	Patient ID	1909221031/AMB
Birthday	03-09-1983	Sample ID	1909221031/AMB
Age at delivery	36.5	Sample Date	16-09-2019
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	65	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	4.88 mIU/ml	1.18	13 + 0
fb-hCG	56.5 ng/ml	1.88	Method
			CRL Robinson
Risks at term			Scan date
Age risk		1:303	14-09-2019
Biochemical T21 risk		1:592	Crown rump length in mm
Combined trisomy 21 risk		1:2399	70.29
Trisomy 13/18 + NT		<1:10000	Nuchal translucency MoM
			0.98
			Nasal bone
			unknown
			Sonographer
			.
			Qualifications in measuring NT
			MD
Risk			Trisomy 21
1:10			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
			After the result of the Trisomy 21 test (with NT) it is expected that among 2399 women with the same data, there is one woman with a trisomy 21 pregnancy and 2398 women with not affected pregnancies. 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			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician