

Patient data			
Name	GURPREET		Patient ID
Birth day	20-05-1989	Sample ID	211902040001
Age at sample date	30.0	Sample Date	04-02-2019
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	unknown
Weight	75	diabetes	unknown
Smoker	unknown	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.8 mIU/ml	0.69	13 + 2
fb-hCG	35.1 ng/ml	1.17	Method
			LMP
			Scan date
			Crown rump length in mm
			Nuchal translucency MoM
			Nasal bone
			Sonographer
			Qualifications in measuring NT
			MD
Risks at sampling date		Trisomy 21	
Age risk	1:790	The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.	
Biochemical T21 risk	1:1460	After the result of the Trisomy 21 test (with NT) it is expected that among 2641 women with the same data, there is one woman with a trisomy 21 pregnancy and 2640 women with not affected pregnancies.	
Combined trisomy 21 risk	1:2641	The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.	
Trisomy 13/18 + NT	<1:10000	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!	
Risk			
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