

Patient data			
Name	MRS. GURPREET KAUR	Patient ID	
Birthday	08-08-1992	Sample ID	211904090002
Age at sample date	26.9	Sample Date	09-04-2019
Gestational age	13 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	58	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	8.62 mIU/ml	1.57	13 + 4
fb-hCG	47.1 ng/ml	1.33	Method
Risks at sampling date			CRL Robinson
Age risk		1:967	Scan date
Biochemical T21 risk		1:7584	09-04-2019
Combined trisomy 21 risk		<1:10000	Crown rump length in mm
Trisomy 13/18 + NT		<1:10000	74.3
			Nuchal translucency MoM
			0.66
			Nasal bone
			present
			Sonographer
			.
			Qualifications in measuring NT
			MD
Risk			Trisomy 21
<p>1:10</p> <p>Age</p>			<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>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><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>			

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Sign of Physician

