

Patient data			
Name	MRS. MANDEEP KAUR	Patient ID	
Birthday	04-02-1994	Sample ID	1905220499/AMB
Age at sample date	25.3	Sample Date	14-05-2019
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	60	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	4.15 mIU/ml	1.00	13 + 0
fb-hCG	35.1 ng/ml	0.92	Method
			CRL Robinson
Risks at sampling date			Scan date
Age risk		1:965	14-05-2019
Biochemical T21 risk		1:7380	Crown rump length in mm
Combined trisomy 21 risk		<1:10000	69
Trisomy 13/18 + NT		<1:10000	Nuchal translucency MoM
			1.27
			Nasal bone
			unknown
			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 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>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician