

Patient data			
Name	MRS. GURINDER (A)	Patient ID	2001220599/AMB (A)
Birthday	07-01-1993	Sample ID	2001220599/AMB (A)
Age at delivery	27.5	Sample Date	17-01-2020
Gestational age	13 + 0		
Correction factors			
Fetuses	2	IVF	no
Weight	41.9	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	10.7 mIU/ml	0.92	12 + 5
fb-hCG	145 ng/ml	1.81	Method
			CRL Robinson
Risks at term		Scan date	15-01-2020
Age risk	1:1205	Crown rump length in mm	65.9
Biochemical T21 risk	1:1554	Nuchal translucency MoM	0.60
Combined trisomy 21 risk	1:8580	Nasal bone	present
Trisomy 13/18 + NT	<1:10000	Sonographer	.
		Qualifications in measuring NT	MD
Risk		Trisomy 21	
1:10		<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 8580 women with the same data, there is one woman with a trisomy 21 pregnancy and 8579 women with not affected pregnancies.</p> <p>The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.</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>			

Sign of Physician