

Patient data			
Name	RS. SUKHJIT KAUR	Patient ID	1707220518/AMB
Birth day	06-07-1985	Sample ID	1707220518/AMB
Age at sample date	32.0	Sample Date	18-07-2017
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	72	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.4 mIU/ml	1.34	12 + 1
fb-hCG	54.7 ng/ml	1.57	Method
Risks at sampling date			CRL Robinson
Age risk		1:477	Scan date
Biochemical T21 risk		1:1870	17-07-2017
Combined trisomy 21 risk		1:8297	Crown rump length in mm
Trisomy 13/18 + NT		<1:10000	58.4
Risk		1:10	Nuchal translucency MoM
			0.92
			Nasal bone
			present
			Sonographer
			.
			Qualifications in measuring NT
			MD
			Trisomy 21
The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.			
<p>After the result of the Trisomy 21 test (with NT) it is expected that among 8297 women with the same data, there is one woman with a trisomy 21 pregnancy and 8296 women with not affected pregnancies.</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>			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician

below cut off

Below Cut Off, but above Age Risk

above cut off

