

Patient data			
Name	MRS. KULVIR	Patient ID	211907030002
Birthday	01/08/1989	Sample ID	211907030002
Age at sample date	29	Sample Date	03/07/19
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	43	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.9 mIU/ml	0.64	12 + 3
fb-hCG	59.6 ng/ml	1.78	Method
			CRL Robinson
			Scan date
			02/07/19
Risks at sampling date			Crown rump length in mm
Age risk		1:959	58.8
Biochemical T21 risk		1:546	Nuchal translucency MoM
Combined trisomy 21 risk		1:3312	0.69
Trisomy 13/18 + NT		<1:10000	Nasal bone
			present
			Sonographer
			Qualifications in measuring NT
			MD
Risk			Trisomy 21
			<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 3312 women with the same data, there is one woman with a trisomy 21 pregnancy and 3311 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 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