

Patient data			
Name	MRS. KARAMJIT	Patient ID	1908221635/AMB
Birthday	25-02-1992	Sample ID	1908221635/AMB
Age at delivery	28.0	Sample Date	27-08-2019
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	61.9	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.33 mIU/ml	0.73	12 + 2
fb-hCG	52.6 ng/ml	1.48	Method
			CRL Robinson
			Scan date
			26-08-2019
Risks at term		Crown rump length in mm	60
Age risk	1:1162	Nuchal translucency MoM	0.83
Biochemical T21 risk	1:1411	Nasal bone	present
Combined trisomy 21 risk	1:7873	Sonographer	.
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT	MD
Risk	1:10	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 7873 women with the same data, there is one woman with a trisomy 21 pregnancy and 7872 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>	
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