

Patient data			
Name	MRS. HARDEEP (B)	Patient ID	1906220663/AMB (B)
Birthday	03-04-1990	Sample ID	1906220663/AMB (B)
Age at sample date	29.2	Sample Date	15-06-2019
Gestational age	12 + 3		
Correction factors			
Fetuses	2	IVF	no
Weight	45	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	10.5 mIU/ml	1.21	Gestational age 12 + 1
fb-hCG	195 ng/ml	1.96	Method CRL Robinson
Risks at sampling date			Scan date 13-06-2019
Age risk		1:706	Crown rump length in mm 58.06
Biochemical T21 risk		1:1314	Nuchal translucency MoM 0.78
Combined trisomy 21 risk		1:6841	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer .
			Qualifications in measuring NT MD
Risk			Trisomy 21
1:10			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 6841 women with the same data, there is one woman with a trisomy 21 pregnancy and 6840 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			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician