

Prisca 5.0.2.37
Date of report: 17-06-2018

Patient data			
Name	MRS. PRIYANKA	Patient ID	1806220392/AMB
Birth day	28-12-1991	Sample ID	1806220392/AMB
Age at sample date	26.5	Sample Date	16-06-2018
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	63	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.34 mIU/ml	0.63	Gestational age 12 + 6
fb-hCG	64.2 ng/ml	1.96	Method CRL Robinson
			Scan date 16-06-2018
Risks at sampling date			Crown rump length in mm 68.7
Age risk		1:900	Nuchal translucency MoM 1.04
Biochemical T21 risk		1:396	Nasal bone present
Combined trisomy 21 risk		1:1545	Sonographer .
Trisomy 13/18 + NT		<1:10000	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 1545 women with the same data, there is one woman with a trisomy 21 pregnancy and 1544 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

below cut off	Below Cut Off, but above Age Risk	above cut off
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