

Prisca 5.0.2.37
Date of report: 22-12-2017

Patient data			
Name	MRS. SHABANAM	Patient ID	1712220342/AMB
Birthdate	11-09-1986	Sample ID	1712220342/AMB
Age at sample date	31.3	Sample Date	21-12-2017
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	59	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.48 mIU/ml	0.97	Gestational age 12 + 3
fb-hCG	24.1 ng/ml	0.68	Method CRL Robinson
Risks at sampling date			Scan date 20-12-2017
Age risk		1:543	Crown rump length in mm 62.2
Biochemical T21 risk		1:7179	Nuchal translucency MoM 1.25
Combined trisomy 21 risk		<1:10000	Nasal bone present
Trisomy 13/18 + NT		<1:10000	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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</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