

KOS DIAGNOSTIC LAB

Patient data			
Name	MRS. RAJNI	Patient ID	1908220038/AMB
Birth day	02-02-1989	Sample ID	1908220038/AMB
Age at delivery	31.0	Sample Date	01-08-2019
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	57	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.7 mIU/ml	0.48	12 + 3
fb-hCG	44.1 ng/ml	1.21	Method
			CRL Robinson
			Scan date
			01-08-2019
Risks at term		Crown rump length in mm	
Age risk	1:851	61.6	
Biochemical T21 risk	1:585	Nuchal translucency MoM	
Combined trisomy 21 risk	1:3624	0.62	
Trisomy 13/18 + NT	<1:10000	Nasal bone	
		present	
		Sonographer	
		Qualifications in measuring NT	
		MD	
Risk		Trisomy 21	
Risk 1:10 		<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 3624 women with the same data, there is one woman with a trisomy 21 pregnancy and 3623 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><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>			

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Sign of Physician

