

KOS DIAGNOSTIC LAB

Patient data			
Name	MRS. JASWINDER KAUR	Patient ID	1909220142/AMB
Birthday	25-11-1986	Sample ID	1909220142/AMB
Age at delivery	33.3	Sample Date	03-09-2019
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	53	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	4.89 mIU/ml	1.20	12 + 3
fb-hCG	64.8 ng/ml	1.77	Method
			CRL Robinson
			Scan date
			02-09-2019
Risks at term		Crown rump length in mm	62
Age risk	1:595	Nuchal translucency MoM	0.87
Biochemical T21 risk	1:1390	Nasal bone	present
Combined trisomy 21 risk	1:6757	Sonographer	.
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT	MD
Risk	1:10	Trisomy 21	
		<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 6757 women with the same data, there is one woman with a trisomy 21 pregnancy and 6756 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

