

KOS DIAGNOSTIC LAB

Patient data			
Name	MRS. PARMINDER	Patient ID	1907221442/AMB
Birthday	20-08-1985	Sample ID	1907221442/AMB
Age at delivery	34.5	Sample Date	26-07-2019
Gestational age	12 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	52	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	7.8 mIU/ml	2.22	12 + 0
fb-hCG	98.3 ng/ml	2.48	Method
			CRL Robinson
			Scan date
			25-07-2019
Risks at term		Crown rump length in mm	
Age risk	1:477	56	
Biochemical T21 risk	1:1167	Nuchal translucency MoM	
Combined trisomy 21 risk	1:5211	0.88	
Trisomy 13/18 + NT	<1:10000	Nasal bone	
		present	
		Sonographer	
		Qualifications in measuring NT	
		MD	
Risk		Trisomy 21	
1:10		<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 5211 women with the same data, there is one woman with a trisomy 21 pregnancy and 5210 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