

KOS DIAGNOSTIC LAB

Patient data			
Name	MRS. KIRAN (B)	Patient ID	
Birthday	03-03-1984	Sample ID	1909221837/AMB
Age at delivery	36.1	Sample Date	30-09-2019
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	60.2	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	10.7 mIU/ml	3.41	Gestational age 12 + 1
fb-hCG	198.4 ng/ml	5.40	Method CRL Robinson
Risks at term			Scan date 29-09-2019
Age risk		1:336	Crown rump length in mm 58.4
Biochemical T21 risk		1:221	Nuchal translucency MoM 0.98
Combined trisomy 21 risk		1:839	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer .
			Qualifications in measuring NT MD
Risk			Trisomy 21
<p>Risk 1:10</p>			<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 839 women with the same data, there is one woman with a trisomy 21 pregnancy and 838 women with not affected pregnancies. The free beta HCG level is high. The PAPP-A level is high. 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! The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)). 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