

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><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 839 women with the same data, there is one woman with a trisomy 21 pregnancy and 838 women with not affected pregnancies.</p> <p>The free beta HCG level is high.</p> <p>The PAPP-A level is high.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.</p> <p>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>			

Sign of Physician

