

KOS DIAGNOSTIC LAB

Patient data			
Name	MRS. CHANDER KANTA	Patient ID	1909220145/AMB
Birth day	02-03-1995	Sample ID	1909220145/AMB
Age at delivery	25.0	Sample Date	03-09-2019
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	50	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	10.5 mIU/ml	2.55	12 + 2
fb-hCG	172 ng/ml	4.50	Method
			CRL Robinson
			Scan date
			02-09-2019
Risks at term		Crown rump length in mm	60
Age risk	1:1377	Nuchal translucency MoM	0.77
Biochemical T21 risk	1:906	Nasal bone	present
Combined trisomy 21 risk	1:4578	Sonographer	.
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT	MD
Risk		Trisomy 21	
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 4578 women with the same data, there is one woman with a trisomy 21 pregnancy and 4577 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><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

