

KOS DIAGNOSTIC LAB

Patient data			
Name	MRS. PARAMJIT	Patient ID	1907221443/AMB
Birthday	11-06-1988	Sample ID	1907221443/AMB
Age at delivery	31.6	Sample Date	26-07-2019
Gestational age	13 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	57	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	4.39 mIU/ml	0.75	13 + 0
fb-hCG	101 ng/ml	3.55	Method
			CRL Robinson
Risks at term			Scan date
Age risk		1:782	20-07-2019
Biochemical T21 risk		1:109	Crown rump length in mm
Combined trisomy 21 risk		1:694	68.8
Trisomy 13/18 + NT		<1:10000	Nuchal translucency MoM
			0.75
			Nasal bone
			present
			Sonographer
			.
			Qualifications in measuring NT
			MD
Risk			Trisomy 21
1:10			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
			After the result of the Trisomy 21 test (with NT) it is expected that among 694 women with the same data, there is one woman with a trisomy 21 pregnancy and 693 women with not affected pregnancies. The free beta HCG 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!
1:100			
1:250			
1:1000			
1:10000			
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			
Age			
Trisomy 13/18 + NT			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician

