

KOS DIAGNOSTIC LAB

Patient data			
Name	MRS. PATWINDER	Patient ID	1907221696/AMB
Birthday	27-09-1991	Sample ID	1907221696/AMB
Age at delivery	28.3	Sample Date	31-07-2019
Gestational age	13 + 4		
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	4.67 mIU/ml	0.88	12 + 5
fb-hCG	67.3 ng/ml	2.25	Method
			CRL Robinson
			Scan date
			25-07-2019
			Crown rump length in mm
			65.71
			Nuchal translucency MoM
			0.85
			Nasal bone
			present
			Sonographer
			.
			Qualifications in measuring NT
			MD
Risks at term		Trisomy 21	
Age risk	1:1133	The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.	
Biochemical T21 risk	1:749	After the result of the Trisomy 21 test (with NT) it is expected that among 4091 women with the same data, there is one woman with a trisomy 21 pregnancy and 4090 women with not affected pregnancies.	
Combined trisomy 21 risk	1:4091	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!	
Trisomy 13/18 + NT	<1:10000	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!	
Risk 1:10			
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

