

KOS DIAGNOSTIC LAB

Patient data			
Name	MRS. SARMILA	Patient ID	
Birthday	19-11-1994	Sample ID	2211220604/AMB
Age at sample date	28.0	Sample Date	25-11-2022
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	52	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	0.7 mIU/ml	0.14	Gestational age
fb-hCG	12.1 ng/ml	0.26	Method
			Scan date
Risks at sampling date			Crown rump length in mm
Age risk		1:799	Nuchal translucency MoM
Biochemical T21 risk		1:510	Nasal bone
Combined trisomy 21 risk		1:3316	Sonographer
Trisomy 13/18 + NT		>1:50	Qualifications in measuring NT
			MD
Trisomy 21			
<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 3316 women with the same data, there is one woman with a trisomy 21 pregnancy and 3315 women with not affected pregnancies.</p> <p>The free beta HCG level is low. The PAPP-A level is low. 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>			
		<p>Trisomy 13/18 + NT</p> <p>The calculated risk for Trisomy 13/18 test (with nuchal translucency) is >1:50, which indicates an increased risk.</p>	
		<p>Sign of Physician</p>	