

KOS DIAGNOSTIC LAB

Patient data			
Name	MRS. AMARJEET	Patient ID	
Birthday	03-08-1994	Sample ID	2209220681/AMB
Age at sample date	28.2	Sample Date	27-09-2022
Gestational age	11 + 5		
Correction factors			
Fetuses	1	IVF	yes
Weight	60.2	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	0.8 mIU/ml	0.29	Gestational age
fb-hCG	114 ng/ml	2.32	Method
			Scan date
Risks at sampling date		Crown rump length in mm	
Age risk	1:761	53.1	
Biochemical T21 risk	>1:50	Nuchal translucency MoM	
Combined trisomy 21 risk	1:191	0.85	
Trisomy 13/18 + NT	<1:10000	Nasal bone	
		present	
		Sonographer	
		Qualifications in measuring NT	
		MD	
Trisomy 21			
<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 191 women with the same data, there is one woman with a trisomy 21 pregnancy and 190 women with not affected pregnancies.</p> <p>The PAPP-A level is low.</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>			

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**Sign of Physician**