

KOS DIAGNOSTIC LAB

Patient data			
Name	MRS. GURJOT KAUR	Patient ID	
Birth day	17-04-1997	Sample ID	2206220204/AMB
Age at sample date	25.1	Sample Date	08-06-2022
Gestational age	11 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	64.9	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.91 mIU/ml	0.82	Gestational age
fb-hCG	151 ng/ml	3.10	Method
			Scan date
Risks at sampling date			Crown rump length in mm
Age risk		1:922	48.98
Biochemical T21 risk		1:230	Nuchal translucency MoM
Combined trisomy 21 risk		1:269	1.36
Trisomy 13/18 + NT		<1:10000	Nasal bone
			present
			Sonographer
			DR. G.B. SINGH
			Qualifications in measuring NT
			M.D
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 269 women with the same data, there is one woman with a trisomy 21 pregnancy and 268 women with not affected pregnancies.</p> <p>The free beta HCG level is high.</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>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician