

KOS DIAG LAB

Patient data			
Name	MRS. MANPREET KAUR	Patient ID	
Birthday	1/11/1988	Sample ID	2409220062/AMB
Age at sample date	35.8	Sample Date	30/08/2024
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	yes
Weight	58	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.21 mIU/ml	0.67	Gestational age
fb-hCG	252 ng/ml	5.77	Method
			CRL Robinson
			Scan date
			28/08/2024
			Crown rump length in mm
			63
			Nuchal translucency MoM
			0.62
			Nasal bone
			present
			Sonographer
			DR. RUPALI GARG
			Qualifications in measuring NT
			M.D
Risks at sampling date			
Age risk		1:229	
Biochemical T21 risk		>1:50	
Combined trisomy 21 risk		1:117	
Trisomy 13/18 + NT		<1:10000	
Risk		Trisomy 21	
		<p>The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 117 women with the same data, there is one woman with a trisomy 21 pregnancy and 116 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 hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>	
		<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>	

Sign of Physician