

KOS DIAG LAB

Patient data			
Name	MRS. KIRAN	Patient ID	
Birthday	25/06/1988	Sample ID	2409220060/AMB
Age at sample date	36.2	Sample Date	1/09/2024
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	59	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.9 mIU/ml	0.88	Gestational age 12 + 3
fb-hCG	115 ng/ml	2.61	Method CRL Robinson
			Scan date 30/08/2024
Risks at sampling date			Crown rump length in mm 61.6
Age risk		1:209	Nuchal translucency MoM 1.00
Biochemical T21 risk		1:96	Nasal bone present
Combined trisomy 21 risk		1:391	Sonographer DR. NIDHI SHARMA
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT M.D
Trisomy 21			
<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 391 women with the same data, there is one woman with a trisomy 21 pregnancy and 390 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>			
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>			

**Sign of Physician**