

Prisca 5.1.0.17  
Date of report: 17/03/25

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Patient data				
Name	Mrs. GULNAZ PARVEEN HAIDAR KHA		Patient ID	0622503120063
Birthday	02/01/99		Sample ID	A1631153
Age at sample date	26.2		Sample Date	12/03/25
Gestational age	12 + 6			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	44	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	0 + 0
PAPP-A	3.87 mIU/mL	0.60	Method	CRL Robinson
fb-hCG	234 ng/mL	5.42	Scan date	11/03/25
Risks at sampling date			Crown rump length in mm	
Age risk	1:915		65.6	
Biochemical T21 risk	1:54		Nuchal translucency MoM	
Combined trisomy 21 risk	1:368		0.42	
Trisomy 13/18 + NT	<1:10000		Nasal bone	
			present	
			Sonographer	
			DR S MANJULA MBBS	
			Qualifications in measuring NT	
			DMCH	
Risk			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 368 women with the same data, there is one woman with a trisomy 21 pregnancy and 367 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. 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>				

Sign of Physician