

Prisca 5.1.0.17
Date of report: 23/12/25

Patient data				
Name	Mrs. SUDHA GRITLAHARE		Patient ID	0622512200066
Birthday	17/07/85		Sample ID	B2602627
Age at sample date	40.4		Sample Date	20/12/25
Gestational age	13 + 3			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	52	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 5
PAPP-A	2.15 mIU/mL	0.33	Method	CRL Robinson
fb-hCG	31.8 ng/mL	0.92	Scan date	15/12/25
Risks at sampling date			Crown rump length in mm	
Age risk	1:73		65	
Biochemical T21 risk	>1:50		Nuchal translucency MoM	
Combined trisomy 21 risk	1:182		0.90	
Trisomy 13/18 + NT	1:1443		Nasal bone	
			present	
			Sonographer	
			NA	
			Qualifications in measuring NT	
			Sonographer	
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 182 women with the same data, there is one woman with a trisomy 21 pregnancy and 181 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. 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 1:1443, which represents a low risk.</p>				

Sign of Physician