

Prisca 5.1.0.17
Date of report: 31-10-2024

N A

Patient data			
Name	Mrs. SAVITA GERNARE		Patient ID
Birthday	16-07-1996	Sample ID	0662410300059
Age at sample date	28.3	Sample Date	A1686559
Gestational age	13 + 0		30-10-2024
Correction factors			
Fetuses	1	IVF	unknown
Weight	56	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies	unknown		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	7.15 mIU/mL	1.38	12 + 6
fb-hCG	50.39 ng/mL	1.32	Method
			CRL Robinson
			Scan date
			29-10-2024
Risks at sampling date		Crown rump length in mm	
Age risk	1:788	67	
Biochemical T21 risk	1:4927	Nuchal translucency MoM	
Combined trisomy 21 risk	1:4869	1.41	
Trisomy 13/18 + NT	<1:10000	Nasal bone	
		present	
		Sonographer	
		N A	
		Qualifications in measuring NT	
		MD	
Risk		Trisomy 21	
		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 4869 women with the same data, there is one woman with a trisomy 21 pregnancy and 4868 women with not affected pregnancies. 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! The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)). The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
Trisomy 13/18 + NT			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician

below cut off
 Below Cut Off, but above Age Risk
 above cut off