

Prisca 5.1.0.17
Date of report: 18/12/24

NA

Patient data			
Name	Mrs. SUNITA GORE	Patient ID	0672412170057
Birthday	10/04/97	Sample ID	A1757687
Age at sample date	27.7	Sample Date	17/12/24
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	39	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.95 mIU/mL	0.50	12 + 1
fb-hCG	40.57 ng/mL	0.78	Method
			CRL Robinson
			Scan date
			16/12/24
Risks at sampling date			Trisomy 21
Age risk	1:809		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
Biochemical T21 risk	1:1643		After the result of the Trisomy 21 test (with NT) it is expected that among 9444 women with the same data, there is one woman with a trisomy 21 pregnancy and 9443 women with not affected pregnancies.
Combined trisomy 21 risk	1:9444		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!
Trisomy 13/18 + NT	<1:10000		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