

Prisca 5.1.0.17
Date of report: 20-12-2024

NA

Patient data			
Name	Mrs. NANDINI NETAM	Patient ID	0622412190032
Birthday	27-03-1993	Sample ID	24241740
Age at sample date	31.7	Sample Date	19-12-2024
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	45	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.75 mIU/mL	0.60	12 + 5
fb-hCG	52.84 ng/mL	1.24	Method
			CRL Robinson
			Scan date
			18-12-2024
Risks at sampling date			Trisomy 21
Age risk	1:512		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
Biochemical T21 risk	1:587		After the result of the Trisomy 21 test (with NT) it is expected that among 2074 women with the same data, there is one woman with a trisomy 21 pregnancy and 2073 women with not affected pregnancies.
Combined trisomy 21 risk	1:2074		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