

Prisca

5.1.0.17

Date of report:

08-05-2023

Patient data			
Name	Mrs. SAPNA	Patient ID	0012305060182
Birthday	03-05-1996	Sample ID	24333532
Age at sample date	27.0	Sample Date	06-05-2023
Gestational age	12 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	48	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.28 mIU/mL	0.52	Gestational age 11 + 6
fb-hCG	43.88 ng/mL	0.89	Method CRL Robinson
			Scan date 04-05-2023
Risks at sampling date			Crown rump length in mm 55
Age risk	1:847		Nuchal translucency MoM 0.69
Biochemical T21 risk	1:1436		Nasal bone present
Combined trisomy 21 risk	1:8330		Sonographer NA
Trisomy 13/18 + NT	<1:10000		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 8330 women with the same data, there is one woman with a trisomy 21 pregnancy and 8329 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