

Prisca

5.1.0.17

Date of report:

23/03/23

Patient data			
Name	Mrs. JAYA SRI		Patient ID
Birthday	18/07/88	Sample ID	0012303210496
Age at sample date	34.7	Sample Date	23940348
Gestational age	11 + 4	Sample Date	21/03/23
Correction factors			
Fetuses	1	IVF	no
Weight	76	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.91 mIU/mL	1.49	11 + 3
fb-hCG	56.32 ng/mL	1.17	Method
			CRL Robinson
			Scan date
			20/03/23
Risks at sampling date			Crown rump length in mm
Age risk			48.9
Biochemical T21 risk			Nuchal translucency MoM
1:281			0.88
Combined trisomy 21 risk			Nasal bone
1:2662			unknown
Trisomy 13/18 + NT			Sonographer
<1:10000			NA
<1:10000			Qualifications in measuring NT
			MD
Risk		Trisomy 21	
1:10		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.	
1:100		After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.	
1:250		The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.	
1:1000		Please note that risk calculations are statistical approaches and have no diagnostic value!	
1:10000		The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).	
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49		The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
Age			
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