

Prisca 5.1.0.17

Date of report: 20-11-2023

Patient data			
Name	Mrs. PUSPHAVATHI	Patient ID	0012311190413
Birthday	19-01-1998	Sample ID	24668761
Age at sample date	25.8	Sample Date	19-11-2023
Gestational age	13 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	63.9	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	23.75 mIU/mL	4.17	Gestational age 13 + 4
fb-hCG	29.22 ng/mL	0.98	Method CRL Robinson
Risks at sampling date			Scan date 18-11-2023
Age risk	1:960		Crown rump length in mm 78.38
Biochemical T21 risk	<1:10000		Nuchal translucency MoM 0.53
Combined trisomy 21 risk	<1:10000		Nasal bone present
Trisomy 13/18 + NT	<1:10000		Sonographer NA
			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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. The PAPP-A level is high. 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.			

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