

Prisca

5.1.0.17

Date of report:

24-09-2021

C.PADMA.MD

Patient data			
Name	Mrs. SWAPNA		Patient ID
Birthday	07-04-1991		Sample ID
Age at sample date	30.5		Sample Date
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	55	diabetes	no
Smoker	no	Origin	Asian
			Previous trisomy 21 pregnancies
			unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.73 mIU/mL	0.61	12 + 4
fb-hCG	45.26 ng/mL	1.06	Method
Risks at sampling date			CRL Robinson
Age risk	1:609		Scan date
Biochemical T21 risk	1:1048		24-09-2021
Combined trisomy 21 risk	1:5807		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		64.62
			Nuchal translucency MoM
			0.85
			Nasal bone
			present
			Sonographer
			DR NA
			Qualifications in measuring NT
			MD
Risk		Trisomy 21	
		<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 5807 women with the same data, there is one woman with a trisomy 21 pregnancy and 5806 women with not affected pregnancies.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>	
Trisomy 13/18 + NT			
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician

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