

Prisca

5.1.0.17

Date of report:

27-08-2024

N A

Patient data			
Name	Mrs. SHANTI SARIN		Patient ID
Birth day	23-03-2005		Sample ID
Age at sample date	19.4		Sample Date
Gestational age	12 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	42	diabetes	no
Smoker	no	Origin	Asian
			Previous trisomy 21 pregnancies
			unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	8.29 mIU/mL	1.63	Method
fb-hCG	44.52 ng/mL	0.86	CRL Robinson
Risks at sampling date			Scan date
Age risk			26-08-2024
Biochemical T21 risk			Crown rump length in mm
Combined trisomy 21 risk			57
Trisomy 13/18 + NT			Nuchal translucency MoM
			0.87
			Nasal bone
			present
			Sonographer
			N A
			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 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