

Prisca

5.1.0.17

Date of report:

02/02/22

Uma maheshwari devi

Patient data			
Name	Mrs. V VAISHNAVI.		Patient ID
Birth day	13/12/99		Sample ID
Age at sample date	22.1		Sample Date
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	52	diabetes	no
Smoker	no	Origin	Asian
			Previous trisomy 21 pregnancies
			unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.99 mIU/mL	0.37	12 + 5
fb-hCG	46.87 ng/mL	1.16	Method
Risks at sampling date			CRL Robinson
Age risk	1:1064		Scan date
Biochemical T21 risk	1:398		31/01/22
Combined trisomy 21 risk	1:2627		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		66
			Nuchal translucency MoM
			0.54
			Nasal bone
			present
			Sonographer
			DR 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 2627 women with the same data, there is one woman with a trisomy 21 pregnancy and 2626 women with not affected pregnancies. The PAPP-A level is low. 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