

Prisca

5.1.0.17

Date of report:

13/02/23

TEJASWI

Patient data			
Name	Mrs. ANISHA		Patient ID
Birthday	21/08/93	Sample ID	0012302110319
Age at sample date	29.5	Sample Date	24171478
Gestational age	12 + 2		11/02/23
Correction factors			
Fetuses	1	IVF	no
Weight	83	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.46 mIU/mL	0.60	12 + 2
fb-hCG	37.25 ng/mL	0.93	Method
			CRL Robinson
			Scan date
			11/02/23
Risks at sampling date			Crown rump length in mm
			60
Age risk	1:681		Nuchal translucency MoM
Biochemical T21 risk	1:1548		0.90
Combined trisomy 21 risk	1:229		Nasal bone
Trisomy 13/18 + NT	<1:10000		absent
			Sonographer
			TEJASWI
			Qualifications in measuring NT
			MD
Risk		Trisomy 21	
		The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk. After the result of the Trisomy 21 test (with NT) it is expected that among 229 women with the same data, there is one woman with a trisomy 21 pregnancy and 228 women with not affected pregnancies. 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