

JITM SKILLS PVT LTD
D-89,SECTOR-2,NOIDA (U.P)

Prisca 5.1.0.17
Date of report: 19/04/23

Patient data			
Name	Mrs.SHRUTI	Patient ID	4243105
Birthday	15/05/92	Sample ID	4243105
Age at delivery	31.4	Sample Date	18/04/23
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	60	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies			unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.85 mIU/ml	0.89	Gestational age 12 + 5
fb-hCG	23.6 ng/ml	0.74	Method CRL Robinson
Risks at term			Scan date 15/04/23
Age risk		1:803	Crown rump length in mm 65.1
Biochemical T21 risk		1:7296	Nuchal translucency MoM 0.78
Combined trisomy 21 risk		<1:10000	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer Dr.Anamika
			Qualifications in measuring NT FMF
			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
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