

JITM DIAGNOSTICS PVT LTD
D – 87 , Sector – 2 , NOIDA (UP) 201301

Prisca 5.1.0.17
Date of report: 11/04/22

Patient data			
Name	Mrs.NISHA 1022846	Patient ID	3673297
Birthday	28/10/97	Sample ID	3673297
Age at delivery	25.0	Sample Date	11/04/22
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	unknown
Weight	61	diabetes	unknown
Smoker	unknown	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.73 mIU/ml	1.01	Gestational age 12 + 5
fb-hCG	30.46 ng/ml	1.02	Method CRL Robinson
Risks at term			Scan date 06/04/22
Age risk		1:1381	Crown rump length in mm 66
Biochemical T21 risk		1:8416	Nuchal translucency MoM 0.71
Combined trisomy 21 risk		<1:10000	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer DR.KRITI RAJ
			Qualifications in measuring NT FMF
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
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