

JITM SKILLS PVT LTD
D-87, SECTOR-2,NOIDA (U.P)

Prisca 5.1.0.17

Date of report: 18/04/23

Patient data			
Name	Mrs.POONAM	Patient ID	4243110
Birth day	05/06/94	Sample ID	4243110
Age at delivery	29.4	Sample Date	17/04/23
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	50	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	5.33 mIU/ml	0.99	12 + 6
fb-hCG	23.6 ng/ml	0.70	Method
			CRL Robinson
			Scan date
			15/04/23
Risks at term			Crown rump length in mm
Age risk	1:1031		67.5
Biochemical T21 risk	<1:10000		Nuchal translucency MoM
Combined trisomy 21 risk	<1:10000		0.58
Trisomy 13/18 + NT	<1:10000		Nasal bone
			present
			Sonographer
			Dr.Anamika
			Qualifications in measuring NT
			FMF
Risk		Trisomy 21	
		<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>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.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>	
Trisomy 13/18 + NT			
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician

